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(54) Title: YIELD-RELATED GENES

(57) Abstract: Recombinant polynucleotides and methods for modifying the phenotype of a plant are provided. In particular, the phenotype that is being modified is a plant's sugar-sensing characteristics.



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YIELD-RELATED GENES

RELATED APPLICATION INFORMATION

The present invention claims the benefit from US Provisional Patent Application Serial
5 Nos. 60/166,228 filed November 17, 1999 and 60/197,899 filed April 17, 2000 and "Plant Trait
Modification III" filed August 22, 2000.

FIELD OF THE INVENTION

This invention relates to the field of plant biology. More particularly, the present
invention pertains to compositions and methods for phenotypically modifying a plant.

BACKGROUND OF THE INVENTION

10 Because sugars are important signaling molecules, the ability to control either the
concentration of a signaling sugar or how the plant perceives or responds to a signaling sugar can
be used to control plant development, physiology or metabolism. For example, the flux of sucrose
(a disaccharide sugar used for systemically transporting carbon and energy in most plants) has
15 been shown to affect gene expression and alter storage compound accumulation in seeds (Wobus
et al (1999) Biol. Chem. 380:937-944). Manipulation of the sucrose signaling pathway in seeds
may therefore cause seeds to have more protein, oil or carbohydrate, depending on the type of
manipulation. Similarly, in tubers, sucrose is converted to starch which is used as an energy store.

It is thought that sugar signaling pathways may partially determine the levels of
20 starch synthesized in the tubers (Zrenner et al. (1996) Plant J. 9:671-681). The manipulation of
sugar signaling in tubers could lead to tubers with a higher starch content. Thus, manipulating the
sugar signal transduction pathway may lead to altered gene expression to produce plants with
desirable traits. In particular, manipulation of sugar signal transduction pathways could be used to
alter source-sink relationships in seeds, tubers, roots and other storage organs leading to an
25 increase in yield.

The present invention provides novel transcription factors useful for modifying a
plant's phenotype in desirable ways by modifying a plant's sugar-sensing characteristics and
thereby, increasing the yield.

SUMMARY OF THE INVENTION

30 In a first aspect, the invention relates to a recombinant polynucleotide comprising
a nucleotide sequence selected from the group consisting of: (a) a nucleotide sequence encoding a
polypeptide comprising a sequence selected from SEQ ID Nos. 2N, where N=1-35, or a
complementary nucleotide sequence thereof; (b) a nucleotide sequence encoding a polypeptide

comprising a conservatively substituted variant of a polypeptide of (a); (c) a nucleotide sequence comprising a sequence selected from those of SEQ ID Nos. 2N-1, where N=1-35, or a complementary nucleotide sequence thereof; (d) a nucleotide sequence comprising silent substitutions in a nucleotide sequence of (c); (e) a nucleotide sequence which hybridizes under
5 stringent conditions over substantially the entire length of a nucleotide sequence of one or more of: (a), (b), (c), or (d); (f) a nucleotide sequence comprising at least 15 consecutive nucleotides of a sequence of any of (a)-(e); (g) a nucleotide sequence comprising a subsequence or fragment of any of (a)-(f), which subsequence or fragment encodes a polypeptide having a biological activity that modifies a plant's sugar-sensing characteristics; (h) a nucleotide sequence having at least
10 34% sequence identity to a nucleotide sequence of any of (a)-(g); (i) a nucleotide sequence having at least 60% identity sequence identity to a nucleotide sequence of any of (a)-(g); (j) a nucleotide sequence which encodes a polypeptide having at least 34% identity sequence identity to a polypeptide of SEQ ID Nos. 2N, where N=1-35; (k) a nucleotide sequence which encodes a polypeptide having at least 60% identity sequence identity to a polypeptide of SEQ ID Nos. 2N,
15 where N=1-35; and (l) a nucleotide sequence which encodes a conserved domain of a polypeptide having at least 65% sequence identity to a conserved domain of a polypeptide of SEQ ID Nos. 2N, where N=1-35. The recombinant polynucleotide may further comprise a constitutive, inducible, or tissue-active promoter operably linked to the nucleotide sequence. The invention also relates to compositions comprising at least two of the above described polynucleotides.

20 In a second aspect, the invention is an isolated or recombinant polypeptide comprising a subsequence of at least about 15 contiguous amino acids encoded by the recombinant or isolated polynucleotide described above.

In another aspect, the invention is a transgenic plant comprising one or more of the above described recombinant polynucleotides. In yet another aspect, the invention is a plant
25 with altered expression levels of a polynucleotide described above or a plant with altered expression or activity levels of an above described polypeptide. Further, the invention is a plant lacking a nucleotide sequence encoding a polypeptide described above. The plant may be a soybean, wheat, corn, potato, cotton, rice, oilseed rape, sunflower, alfalfa, sugarcane, turf, banana, blackberry, blueberry, strawberry, raspberry, cantaloupe, carrot, cauliflower, coffee,
30 cucumber, eggplant, grapes, honeydew, lettuce, mango, melon, onion, papaya, peas, peppers, pineapple, spinach, squash, sweet corn, tobacco, tomato, watermelon, rosaceous fruits, or vegetable brassicas plant.

In a further aspect, the invention relates to a cloning or expression vector comprising the isolated or recombinant polynucleotide described above or cells comprising the cloning or expression vector.

5 In yet a further aspect, the invention relates to a composition produced by incubating a polynucleotide of the invention with a nuclease, a restriction enzyme, a polymerase; a polymerase and a primer; a cloning vector, or with a cell.

10 Furthermore, the invention relates to a method for producing a plant having improved sugar-sensing traits. The method comprises altering the expression of an isolated or recombinant polynucleotide of the invention or altering the expression or activity of a polypeptide of the invention in a plant to produce a modified plant, and selecting the modified plant for modified sugar-sensing traits.

15 In another aspect, the invention relates to a method of identifying a factor that is modulated by or interacts with a polypeptide encoded by a polynucleotide of the invention. The method comprises expressing a polypeptide encoded by the polynucleotide in a plant; and identifying at least one factor that is modulated by or interacts with the polypeptide. In one embodiment the method for identifying modulating or interacting factors is by detecting binding by the polypeptide to a promoter sequence, or by detecting interactions between an additional protein and the polypeptide in a yeast two hybrid system, or by detecting expression of a factor by hybridization to a microarray, subtractive hybridization or differential display.

20 In yet another aspect, the invention is a method of identifying a molecule that modulates activity or expression of a polynucleotide or polypeptide of interest. The method comprises placing the molecule in contact with a plant comprising the polynucleotide or polypeptide encoded by the polynucleotide of the invention and monitoring one or more of the expression level of the polynucleotide in the plant, the expression level of the polypeptide in the plant, and modulation of an activity of the polypeptide in the plant.

25 In yet another aspect, the invention relates to an integrated system, computer or computer readable medium comprising one or more character strings corresponding to a polynucleotide of the invention, or to a polypeptide encoded by the polynucleotide. The integrated system, computer or computer readable medium may comprise a link between one or more sequence strings to a modified plant sugar-sensing trait.

30 In yet another aspect, the invention is a method for identifying a sequence similar or homologous to one or more polynucleotides of the invention, or one or more polypeptides encoded by the polynucleotides. The method comprises providing a sequence database; and, querying the sequence database with one or more target sequences corresponding to the one or

more polynucleotides or to the one or more polypeptides to identify one or more sequence members of the database that display sequence similarity or homology to one or more of the one or more target sequences.

The method may further comprise of linking the one or more of the
5 polynucleotides of the invention, or encoded polypeptides, to a modified plant sugar-sensing phenotype.

BRIEF DESCRIPTION OF THE DRAWINGS

Figure 1 provides a table of exemplary polynucleotide and polypeptide sequences of the invention. The table includes from left to right for each sequence: the SEQ ID No., the
10 internal code reference number (GID), whether the sequence is a polynucleotide or polypeptide sequence, and identification of any conserved domains for the polypeptide sequences.

Figure 2 provides a table of exemplary sequences that are homologous to other sequences provided in the Sequence Listing and that are derived from *Arabidopsis thaliana*. The table includes from left to right: the SEQ ID No., the internal code reference number (GID),
15 identification of the homologous sequence, whether the sequence is a polynucleotide or polypeptide sequence, and identification of any conserved domains for the polypeptide sequences.

Figure 3 provides a table of exemplary sequences that are homologous to the sequences provided in Figures 1 and 2 and that are derived from plants other than *Arabidopsis thaliana*. The table includes from left to right: the SEQ ID No., the internal code reference
20 number (GID), the unique GenBank sequence ID No. (NID), the probability that the comparison was generated by chance (P-value), and the species from which the homologous gene was identified.

25 DETAILED DESCRIPTION

The present invention relates to polynucleotides and polypeptides, e.g. for modifying phenotypes of plants.

In particular, the polynucleotides or polypeptides are useful for modifying traits associated with a plant's sugar-sensing characteristics when the expression levels of the
30 polynucleotides or expression levels or activity levels of the polypeptides are altered. Sugars are central regulatory molecules that control aspects of physiology, metabolism and development. Therefore, the polynucleotides and polypeptides are useful for modifying the growth and germination rates of plants, photosynthesis, glyoxylate metabolism, respiration, starch and

sucrose synthesis and degradation, pathogen response, wounding response, cell cycle regulation, pigmentation, flowering and senescence of plants and for modifying sink-source relationships in seeds, tubers, roots and other storage organs leading to an increase in yield.

The polynucleotides of the invention encode plant transcription factors. The plant transcription factors are derived, e.g., from *Arabidopsis thaliana* and can belong, e.g., to one or more of the following transcription factor families: the AP2 (APETALA2) domain transcription factor family (Riechmann and Meyerowitz (1998) J. Biol. Chem. 379:633-646); the MYB transcription factor family (Martin and Paz-Ares (1997) Trends Genet. 13:67-73); the MADS domain transcription factor family (Riechmann and Meyerowitz (1997) J. Biol. Chem. 378:1079-1101); the WRKY protein family (Ishiguro and Nakamura (1994) Mol. Gen. Genet. 244:563-571); the ankyrin-repeat protein family (Zhang et al. (1992) Plant Cell 4:1575-1588); the miscellaneous protein (MISC) family (Kim et al. (1997) Plant J. 11:1237-1251); the zinc finger protein (Z) family (Klug and Schwabe (1995) FASEB J. 9: 597-604); the homeobox (HB) protein family (Duboule (1994) Guidebook to the Homeobox Genes, Oxford University Press); the CAAT-element binding proteins (Forsburg and Guarente (1989) Genes Dev. 3:1166-1178); the squamosa promoter binding proteins (SPB) (Klein et al. (1996) Mol. Gen. Genet. 1996 250:7-16); the NAM protein family; the IAA/AUX proteins (Rouse et al. (1998) Science 279:1371-1373); the HLH/MYC protein family (Littlewood et al. (1994) Prot. Profile 1:639-709); the DNA-binding protein (DBP) family (Tucker et al. (1994) EMBO J. 13:2994-3002); the bZIP family of transcription factors (Foster et al. (1994) FASEB J. 8:192-200); the BPF-1 protein (Box P-binding factor) family (da Costa e Silva et al. (1993) Plant J. 4:125-135); and the golden protein (GLD) family (Hall et al. (1998) Plant Cell 10:925-936).

In addition to methods for modifying a plant phenotype by employing one or more polynucleotides and polypeptides of the invention described herein, the polynucleotides and polypeptides of the invention have a variety of additional uses. These uses include their use in the recombinant production (i.e., expression) of proteins; as regulators of plant gene expression, as diagnostic probes for the presence of complementary or partially complementary nucleic acids (including for detection of natural coding nucleic acids); as substrates for further reactions, e.g., mutation reactions, PCR reactions, or the like, or as substrates for cloning e.g., including digestion or ligation reactions, and for identifying exogenous or endogenous modulators of the transcription factors.

DEFINITIONS

A “polynucleotide” is a nucleic acid sequence comprising a plurality of polymerized nucleotide residues, e.g., at least about 15 consecutive polymerized nucleotide residues, optionally at least about 30 consecutive nucleotides, at least about 50 consecutive nucleotides. In many instances, a polynucleotide comprises a nucleotide sequence encoding a polypeptide (or protein) or a domain or fragment thereof. Additionally, the polynucleotide may comprise a promoter, an intron, an enhancer region, a polyadenylation site, a translation initiation site, 5' or 3' untranslated regions, a reporter gene, a selectable marker, or the like. The polynucleotide can be single stranded or double stranded DNA or RNA. The polynucleotide optionally comprises modified bases or a modified backbone. The polynucleotide can be, e.g., genomic DNA or RNA, a transcript (such as an mRNA), a cDNA, a PCR product, a cloned DNA, a synthetic DNA or RNA, or the like. The polynucleotide can comprise a sequence in either sense or antisense orientations.

A “recombinant polynucleotide” is a polynucleotide that is not in its native state, e.g., the polynucleotide comprises a nucleotide sequence not found in nature, or the polynucleotide is in a context other than that in which it is naturally found, e.g., separated from nucleotide sequences with which it typically is in proximity in nature, or adjacent (or contiguous with) nucleotide sequences with which it typically is not in proximity. For example, the sequence at issue can be cloned into a vector, or otherwise recombined with one or more additional nucleic acid.

An “isolated polynucleotide” is a polynucleotide whether naturally occurring or recombinant, that is present outside the cell in which it is typically found in nature, whether purified or not. Optionally, an isolated polynucleotide is subject to one or more enrichment or purification procedures, e.g., cell lysis, extraction, centrifugation, precipitation, or the like.

A “recombinant polypeptide” is a polypeptide produced by translation of a recombinant polynucleotide. An “isolated polypeptide,” whether a naturally occurring or a recombinant polypeptide, is more enriched in (or out of) a cell than the polypeptide in its natural state in a wild type cell, e.g., more than about 5% enriched, more than about 10% enriched, or more than about 20%, or more than about 50%, or more, enriched, i.e., alternatively denoted: 105%, 110%, 120%, 150% or more, enriched relative to wild type standardized at 100%. Such an enrichment is not the result of a natural response of a wild type plant. Alternatively, or additionally, the isolated polypeptide is separated from other cellular components with which it is typically associated, e.g., by any of the various protein purification methods herein.

The term "transgenic plant" refers to a plant that contains genetic material, not found in a wild type plant of the same species, variety or cultivar. The genetic material may include a transgene, an insertional mutagenesis event (such as by transposon or T-DNA insertional mutagenesis), an activation tagging sequence, a mutated sequence, a homologous recombination event or a sequence modified by chimeraplasty. Typically, the foreign genetic material has been introduced into the plant by human manipulation.

A transgenic plant may contain an expression vector or cassette. The expression cassette typically comprises a polypeptide-encoding sequence operably linked (i.e., under regulatory control of) to appropriate inducible or constitutive regulatory sequences that allow for the expression of polypeptide. The expression cassette can be introduced into a plant by transformation or by breeding after transformation of a parent plant. A plant refers to a whole plant as well as to a plant part, such as seed, fruit, leaf, or root, plant tissue, plant cells or any other plant material, e.g., a plant explant, as well as to progeny thereof, and to *in vitro* systems that mimic biochemical or cellular components or processes in a cell.

The phrase "ectopically expression or altered expression" in reference to a polynucleotide indicates that the pattern of expression in, e.g., a transgenic plant or plant tissue, is different from the expression pattern in a wild type plant or a reference plant of the same species. For example, the polynucleotide or polypeptide is expressed in a cell or tissue type other than a cell or tissue type in which the sequence is expressed in the wild type plant, or by expression at a time other than at the time the sequence is expressed in the wild type plant, or by a response to different inducible agents, such as hormones or environmental signals, or at different expression levels (either higher or lower) compared with those found in a wild type plant. The term also refers to altered expression patterns that are produced by lowering the levels of expression to below the detection level or completely abolishing expression. The resulting expression pattern can be transient or stable, constitutive or inducible. In reference to a polypeptide, the term "ectopic expression or altered expression" further may relate to altered activity levels resulting from the interactions of the polypeptides with exogenous or endogenous modulators or from interactions with factors or as a result of the chemical modification of the polypeptides.

The term "fragment" or "domain," with respect to a polypeptide, refers to a subsequence of the polypeptide. In some cases, the fragment or domain, is a subsequence of the polypeptide which performs at least one biological function of the intact polypeptide in substantially the same manner, or to a similar extent, as does the intact polypeptide. For example, a polypeptide fragment can comprise a recognizable structural motif or functional domain such as a DNA binding domain that binds to a DNA promoter region, an activation domain or a domain

for protein-protein interactions. Fragments can vary in size from as few as 6 amino acids to the full length of the intact polypeptide, but are preferably at least about 30 amino acids in length and more preferably at least about 60 amino acids in length. In reference to a nucleotide sequence, “a fragment” refers to any subsequence of a polynucleotide, typically, of at least consecutive about
5 15 nucleotides, preferably at least about 30 nucleotides, more preferably at least about 50, of any of the sequences provided herein.

The term “trait” refers to a physiological, morphological, biochemical or physical characteristic of a plant or particular plant material or cell. In some instances, this characteristic is visible to the human eye, such as seed or plant size, or can be measured by available
10 biochemical techniques, such as the protein, starch or oil content of seed or leaves or by the observation of the expression level of genes, e.g., by employing Northern analysis, RT-PCR, microarray gene expression assays or reporter gene expression systems, or by agricultural observations such as stress tolerance, yield or pathogen tolerance.

“Trait modification” refers to a detectable difference in a characteristic in a plant
15 ectopically expressing a polynucleotide or polypeptide of the present invention relative to a plant not doing so, such as a wild type plant. In some cases, the trait modification can be evaluated quantitatively. For example, the trait modification can entail at least about a 2% increase or decrease in an observed trait (difference), at least a 5% difference, at least about a 10% difference, at least about a 20% difference, at least about a 30%, at least about a 50%, at least
20 about a 70%, or at least about a 100%, or an even greater difference. It is known that there can be a natural variation in the modified trait. Therefore, the trait modification observed entails a change of the normal distribution of the trait in the plants compared with the distribution observed in wild type plant.

Trait modifications of particular interest include those to seed (such as embryo
25 or endosperm), fruit, root, flower, leaf, stem, shoot, seedling or the like, including: enhanced tolerance to environmental conditions including freezing, chilling, heat, drought, water saturation, radiation and ozone; improved tolerance to microbial, fungal or viral diseases; improved tolerance to pest infestations, including nematodes, mollicutes, parasitic higher plants or the like; decreased herbicide sensitivity; improved tolerance of heavy metals or enhanced ability to take up
30 heavy metals; improved growth under poor photoconditions (e.g., low light and/or short day length), or changes in expression levels of genes of interest. Other phenotype that can be modified relate to the production of plant metabolites, such as variations in the production of taxol, tocopherol, tocotrienol, sterols, phytosterols, vitamins, wax monomers, anti-oxidants, amino acids, lignins, cellulose, tannins, prenillipids (such as chlorophylls and carotenoids),

glucosinolates, and terpenoids, enhanced or compositionally altered protein or oil production (especially in seeds), or modified sugar (insoluble or soluble) and/or starch composition.

Physical plant characteristics that can be modified include cell development (such as the number of trichomes), fruit and seed size and number, yields of plant parts such as stems, leaves and roots, the stability of the seeds during storage, characteristics of the seed pod (e.g., susceptibility to shattering), root hair length and quantity, internode distances, or the quality of seed coat. Plant growth characteristics that can be modified include growth rate, germination rate of seeds, vigor of plants and seedlings, leaf and flower senescence, male sterility, apomixis, flowering time, flower abscission, rate of nitrogen uptake, biomass or transpiration characteristics, as well as plant architecture characteristics such as apical dominance, branching patterns, number of organs, organ identity, organ shape or size.

POLYPEPTIDES AND POLYNUCLEOTIDES OF THE INVENTION

The present invention provides, among other things, transcription factors (TFs), and transcription factor homologue polypeptides, and isolated or recombinant polynucleotides encoding the polypeptides. These polypeptides and polynucleotides may be employed to modify a plant's sugar-sensing characteristics..

Exemplary polynucleotides encoding the polypeptides of the invention were identified in the *Arabidopsis thaliana* GenBank database using publicly available sequence analysis programs and parameters. Sequences initially identified were then further characterized to identify sequences comprising specified sequence strings corresponding to sequence motifs present in families of known transcription factors. Polynucleotide sequences meeting such criteria were confirmed as transcription factors.

Additional polynucleotides of the invention were identified by screening *Arabidopsis thaliana* and/or other plant cDNA libraries with probes corresponding to known transcription factors under low stringency hybridization conditions. Additional sequences, including full length coding sequences were subsequently recovered by the rapid amplification of cDNA ends (RACE) procedure, using a commercially available kit according to the manufacturer's instructions. Where necessary, multiple rounds of RACE are performed to isolate 5' and 3' ends. The full length cDNA was then recovered by a routine end-to-end polymerase chain reaction (PCR) using primers specific to the isolated 5' and 3' ends. Exemplary sequences are provided in the Sequence Listing.

The polynucleotides of the invention were ectopically expressed in overexpressor or knockout plants and changes in the sugar-sensing characteristics of the plants were observed.

Therefore, the polynucleotides and polypeptides can be employed to improve the sugar-sensing characteristics of plants.

Making polynucleotides

The polynucleotides of the invention include sequences that encode transcription factors and transcription factor homologue polypeptides and sequences complementary thereto, as well as unique fragments of coding sequence, or sequence complementary thereto. Such polynucleotides can be, e.g., DNA or RNA, e.g., mRNA, cRNA, synthetic RNA, genomic DNA, cDNA synthetic DNA, oligonucleotides, etc. The polynucleotides are either double-stranded or single-stranded, and include either, or both sense (i.e., coding) sequences and antisense (i.e., non-coding, complementary) sequences. The polynucleotides include the coding sequence of a transcription factor, or transcription factor homologue polypeptide, in isolation, in combination with additional coding sequences (e.g., a purification tag, a localization signal, as a fusion-protein, as a pre-protein, or the like), in combination with non-coding sequences (e.g., introns or inteins, regulatory elements such as promoters, enhancers, terminators, and the like), and/or in a vector or host environment in which the polynucleotide encoding a transcription factor or transcription factor homologue polypeptide is an endogenous or exogenous gene.

A variety of methods exist for producing the polynucleotides of the invention. Procedures for identifying and isolating DNA clones are well known to those of skill in the art, and are described in, e.g., Berger and Kimmel, Guide to Molecular Cloning Techniques, Methods in Enzymology volume 152 Academic Press, Inc., San Diego, CA ("Berger"); Sambrook et al., Molecular Cloning - A Laboratory Manual (2nd Ed.), Vol. 1-3, Cold Spring Harbor Laboratory, Cold Spring Harbor, New York, 1989 ("Sambrook") and Current Protocols in Molecular Biology, F.M. Ausubel et al., eds., Current Protocols, a joint venture between Greene Publishing Associates, Inc. and John Wiley & Sons, Inc., (supplemented through 2000) ("Ausubel").

Alternatively, polynucleotides of the invention, can be produced by a variety of in vitro amplification methods adapted to the present invention by appropriate selection of specific or degenerate primers. Examples of protocols sufficient to direct persons of skill through in vitro amplification methods, including the polymerase chain reaction (PCR) the ligase chain reaction (LCR), Qbeta-replicase amplification and other RNA polymerase mediated techniques (e.g., NASBA), e.g., for the production of the homologous nucleic acids of the invention are found in Berger, Sambrook, and Ausubel, as well as Mullis et al., (1987) PCR Protocols A Guide to Methods and Applications (Innis et al. eds) Academic Press Inc. San Diego, CA (1990) (Innis). Improved methods for cloning in vitro amplified nucleic acids are described in Wallace et al., U.S. Pat. No. 5,426,039. Improved methods for amplifying large nucleic acids by PCR are

summarized in Cheng et al. (1994) Nature 369: 684-685 and the references cited therein, in which PCR amplicons of up to 40kb are generated. One of skill will appreciate that essentially any RNA can be converted into a double stranded DNA suitable for restriction digestion, PCR expansion and sequencing using reverse transcriptase and a polymerase. See, e.g., Ausubel, Sambrook and Berger, *all supra*.

Alternatively, polynucleotides and oligonucleotides of the invention can be assembled from fragments produced by solid-phase synthesis methods. Typically, fragments of up to approximately 100 bases are individually synthesized and then enzymatically or chemically ligated to produce a desired sequence, e.g., a polynucleotide encoding all or part of a transcription factor. For example, chemical synthesis using the phosphoramidite method is described, e.g., by Beaucage et al. (1981) Tetrahedron Letters 22:1859-69; and Matthes et al. (1984) EMBO J. 3:801-5. According to such methods, oligonucleotides are synthesized, purified, annealed to their complementary strand, ligated and then optionally cloned into suitable vectors. And if so desired, the polynucleotides and polypeptides of the invention can be custom ordered from any of a number of commercial suppliers.

HOMOLOGOUS SEQUENCES

Sequences homologous, i.e., that share significant sequence identity or similarity, to those provided in the Sequence Listing, derived from *Arabidopsis thaliana* or from other plants of choice are also an aspect of the invention. Homologous sequences can be derived from any plant including monocots and dicots and in particular agriculturally important plant species, including but not limited to, crops such as soybean, wheat, corn, potato, cotton, rice, oilseed rape (including canola), sunflower, alfalfa, sugarcane and turf; or fruits and vegetables, such as banana, blackberry, blueberry, strawberry, and raspberry, cantaloupe, carrot, cauliflower, coffee, cucumber, eggplant, grapes, honeydew, lettuce, mango, melon, onion, papaya, peas, peppers, pineapple, spinach, squash, sweet corn, tobacco, tomato, watermelon, rosaceous fruits (such as apple, peach, pear, cherry and plum) and vegetable brassicas (such as broccoli, cabbage, cauliflower, brussel sprouts and kohlrabi). Other crops, fruits and vegetables whose phenotype can be changed include barley, rye, millet, sorghum, currant, avocado, citrus fruits such as oranges, lemons, grapefruit and tangerines, artichoke, cherries, nuts such as the walnut and peanut, endive, leek, roots, such as arrowroot, beet, cassava, turnip, radish, yam, and sweet potato, and beans. The homologous sequences may also be derived from woody species, such as pine, poplar and eucalyptus.

Transcription factors that are homologous to the listed sequences will typically share at least about 34% amino acid sequence identity. More closely related transcription factors can share at least about 50%, about 60%, about 65%, about 70%, about 75% or about 80% or about 90% or about 95% or about 98% or more sequence identity with the listed sequences.

- 5 Factors that are most closely related to the listed sequences share, e.g., at least about 85%, about 90% or about 95% or more % sequence identity to the listed sequences. At the nucleotide level, the sequences will typically share at least about 40% nucleotide sequence identity, preferably at least about 50%, about 60%, about 70% or about 80% sequence identity, and more preferably about 85%, about 90%, about 95% or about 97% or more sequence identity to one or more of the
- 10 listed sequences. The degeneracy of the genetic code enables major variations in the nucleotide sequence of a polynucleotide while maintaining the amino acid sequence of the encoded protein. Conserved domains within a transcription factor family may exhibit a higher degree of sequence homology, such as at least 65% sequence identity including conservative substitutions, and preferably at least 80% sequence identity.

- 15 Identifying Nucleic Acids by Hybridization
Polynucleotides homologous to the sequences illustrated in the Sequence Listing can be identified, e.g., by hybridization to each other under stringent or under highly stringent conditions. Single stranded polynucleotides hybridize when they associate based on a variety of well characterized physico-chemical forces, such as hydrogen bonding, solvent exclusion, base
- 20 stacking and the like. The stringency of a hybridization reflects the degree of sequence identity of the nucleic acids involved, such that the higher the stringency, the more similar are the two polynucleotide strands. Stringency is influenced by a variety of factors, including temperature, salt concentration and composition, organic and non-organic additives, solvents, etc. present in both the hybridization and wash solutions and incubations (and number), as described in more
- 25 detail in the references cited above.

- An example of stringent hybridization conditions for hybridization of complementary nucleic acids which have more than 100 complementary residues on a filter in a Southern or northern blot is about 5°C to 20°C lower than the thermal melting point (T_m) for the specific sequence at a defined ionic strength and pH. The T_m is the temperature (under defined
- 30 ionic strength and pH) at which 50% of the target sequence hybridizes to a perfectly matched probe. Nucleic acid molecules that hybridize under stringent conditions will typically hybridize to a probe based on either the entire cDNA or selected portions, e.g., to a unique subsequence, of the cDNA under wash conditions of 0.2x SSC to 2.0 x SSC, 0.1% SDS at 50-65° C, for example 0.2 x SSC, 0.1% SDS at 65° C. For identification of less closely related homologues washes can

be performed at a lower temperature, e.g., 50° C. In general, stringency is increased by raising the wash temperature and/or decreasing the concentration of SSC.

As another example, stringent conditions can be selected such that an oligonucleotide that is perfectly complementary to the coding oligonucleotide hybridizes to the coding oligonucleotide with at least about a 5-10x higher signal to noise ratio than the ratio for hybridization of the perfectly complementary oligonucleotide to a nucleic acid encoding a transcription factor known as of the filing date of the application. Conditions can be selected such that a higher signal to noise ratio is observed in the particular assay which is used, e.g., about 15x, 25x, 35x, 50x or more. Accordingly, the subject nucleic acid hybridizes to the unique coding oligonucleotide with at least a 2x higher signal to noise ratio as compared to hybridization of the coding oligonucleotide to a nucleic acid encoding known polypeptide. Again, higher signal to noise ratios can be selected, e.g., about 5x, 10x, 25x, 35x, 50x or more. The particular signal will depend on the label used in the relevant assay, e.g., a fluorescent label, a colorimetric label, a radio active label, or the like.

Alternatively, transcription factor homologue polypeptides can be obtained by screening an expression library using antibodies specific for one or more transcription factors. With the provision herein of the disclosed transcription factor, and transcription factor homologue nucleic acid sequences, the encoded polypeptide(s) can be expressed and purified in a heterologous expression system (e.g., *E. coli*) and used to raise antibodies (monoclonal or polyclonal) specific for the polypeptide(s) in question. Antibodies can also be raised against synthetic peptides derived from transcription factor, or transcription factor homologue, amino acid sequences. Methods of raising antibodies are well known in the art and are described in Harlow and Lane (1988) Antibodies: A Laboratory Manual, Cold Spring Harbor Laboratory, New York. Such antibodies can then be used to screen an expression library produced from the plant from which it is desired to clone additional transcription factor homologues, using the methods described above. The selected cDNAs can be confirmed by sequencing and enzymatic activity.

SEQUENCE VARIATIONS

It will readily be appreciated by those of skill in the art, that any of a variety of polynucleotide sequences are capable of encoding the transcription factors and transcription factor homologue polypeptides of the invention. Due to the degeneracy of the genetic code, many different polynucleotides can encode identical and/or substantially similar polypeptides in addition to those sequences illustrated in the Sequence Listing.

For example, Table 1 illustrates, e.g., that the codons AGC, AGT, TCA, TCC, TCG, and TCT all encode the same amino acid: serine. Accordingly, at each position in the sequence where there is a codon encoding serine, any of the above trinucleotide sequences can be used without altering the encoded polypeptide.

5

Table 1

Amino acids			Codon							
Alanine	Ala	A	GCA	GCC	GCG	GCU				
Cysteine	Cys	C	TGC	TGT						
Aspartic acid	Asp	D	GAC	GAT						
Glutamic acid	Glu	E	GAA	GAG						
Phenylalanine	Phe	F	TTC	TTT						
Glycine	Gly	G	GGA	GGC	GGG	GGT				
Histidine	His	H	CAC	CAT						
Isoleucine	Ile	I	ATA	ATC	ATT					
Lysine	Lys	K	AAA	AAG						
Leucine	Leu	L	TTA	TTG	CTA	CTC	CTG	CTT		
Methionine	Met	M	ATG							
Asparagine	Asn	N	AAC	AAT						
Proline	Pro	P	CCA	CCC	CCG	CCT				
Glutamine	Gln	Q	CAA	CAG						
Arginine	Arg	R	AGA	AGG	CGA	CGC	CGG	CGT		
Serine	Ser	S	AGC	AGT	TCA	TCC	TCG	TCT		
Threonine	Thr	T	ACA	ACC	ACG	ACT				
Valine	Val	V	GTA	GTC	GTG	GTT				
Tryptophan	Trp	W	TGG							
Tyrosine	Tyr	Y	TAC	TAT						

Sequence alterations that do not change the amino acid sequence encoded by the polynucleotide are termed "silent" variations. With the exception of the codons ATG and TGG, encoding methionine and tryptophan, respectively, any of the possible codons for the same amino acid can be substituted by a variety of techniques, e.g., site-directed mutagenesis, available in the art. Accordingly, any and all such variations of a sequence selected from the above table are a feature of the invention.

In addition to silent variations, other conservative variations that alter one, or a few amino acids in the encoded polypeptide, can be made without altering the function of the polypeptide, these conservative variants are, likewise, a feature of the invention.

For example, substitutions, deletions and insertions introduced into the sequences provided in the Sequence Listing are also envisioned by the invention. Such sequence modifications can be engineered into a sequence by site-directed mutagenesis (Wu (ed.) Meth. Enzymol. (1993) vol. 217, Academic Press) or the other methods noted below. Amino acid

substitutions are typically of single residues; insertions usually will be on the order of about from 1 to 10 amino acid residues; and deletions will range about from 1 to 30 residues. In preferred embodiments, deletions or insertions are made in adjacent pairs, e.g., a deletion of two residues or insertion of two residues. Substitutions, deletions, insertions or any combination thereof can be combined to arrive at a sequence. The mutations that are made in the polynucleotide encoding the transcription factor should not place the sequence out of reading frame and should not create complementary regions that could produce secondary mRNA structure. Preferably, the polypeptide encoded by the DNA performs the desired function.

Conservative substitutions are those in which at least one residue in the amino acid sequence has been removed and a different residue inserted in its place. Such substitutions generally are made in accordance with the Table 2 when it is desired to maintain the activity of the protein. Table 2 shows amino acids which can be substituted for an amino acid in a protein and which are typically regarded as conservative substitutions.

Table 2

Residue	Conservative Substitutions
Ala	Ser
Arg	Lys
Asn	Gln; His
Asp	Glu
Gln	Asn
Cys	Ser
Glu	Asp
Gly	Pro
His	Asn; Gln
Ile	Leu, Val
Leu	Ile; Val
Lys	Arg; Gln
Met	Leu; Ile
Phe	Met; Leu; Tyr
Ser	Thr; Gly
Thr	Ser; Val
Trp	Tyr
Tyr	Trp; Phe
Val	Ile; Leu

- Substitutions that are less conservative than those in Table 2 can be selected by picking residues that differ more significantly in their effect on maintaining (a) the structure of the polypeptide backbone in the area of the substitution, for example, as a sheet or helical conformation, (b) the charge or hydrophobicity of the molecule at the target site, or (c) the bulk of the side chain. The substitutions which in general are expected to produce the greatest changes in protein properties will be those in which (a) a hydrophilic residue, e.g., seryl or threonyl, is substituted for (or by) a hydrophobic residue, e.g., leucyl, isoleucyl, phenylalanyl, valyl or alanyl; (b) a cysteine or proline is substituted for (or by) any other residue; (c) a residue having an electropositive side chain, e.g., lysyl, arginyl, or histidyl, is substituted for (or by) an electronegative residue, e.g., glutamyl or aspartyl; or (d) a residue having a bulky side chain, e.g., phenylalanine, is substituted for (or by) one not having a side chain, e.g., glycine.

FURTHER MODIFYING SEQUENCES OF THE INVENTION—MUTATION/ FORCED EVOLUTION

In addition to generating silent or conservative substitutions as noted, above, the present invention optionally includes methods of modifying the sequences of the Sequence Listing. In the methods, nucleic acid or protein modification methods are used to alter the given sequences to produce new sequences and/or to chemically or enzymatically modify given sequences to change the properties of the nucleic acids or proteins.

Thus, in one embodiment, given nucleic acid sequences are modified, e.g., according to standard mutagenesis or artificial evolution methods to produce modified sequences. For example, Ausubel, *supra*, provides additional details on mutagenesis methods. Artificial forced evolution methods are described, e.g., by Stemmer (1994) *Nature* 370:389-391, and Stemmer (1994) *Proc. Natl. Acad. Sci. USA* 91:10747-10751. Many other mutation and evolution methods are also available and expected to be within the skill of the practitioner.

Similarly, chemical or enzymatic alteration of expressed nucleic acids and polypeptides can be performed by standard methods. For example, sequence can be modified by addition of lipids, sugars, peptides, organic or inorganic compounds, by the inclusion of modified nucleotides or amino acids, or the like. For example, protein modification techniques are illustrated in Ausubel, *supra*. Further details on chemical and enzymatic modifications can be found herein. These modification methods can be used to modify any given sequence, or to modify any sequence produced by the various mutation and artificial evolution modification methods noted herein.

Accordingly, the invention provides for modification of any given nucleic acid by mutation, evolution, chemical or enzymatic modification, or other available methods, as well as for the products produced by practicing such methods, e.g., using the sequences herein as a starting substrate for the various modification approaches.

For example, optimized coding sequence containing codons preferred by a particular prokaryotic or eukaryotic host can be used e.g., to increase the rate of translation or to produce recombinant RNA transcripts having desirable properties, such as a longer half-life, as compared with transcripts produced using a non-optimized sequence. Translation stop codons can also be modified to reflect host preference. For example, preferred stop codons for *S. cerevisiae* and mammals are TAA and TGA, respectively. The preferred stop codon for monocotyledonous plants is TGA, whereas insects and *E. coli* prefer to use TAA as the stop codon.

The polynucleotide sequences of the present invention can also be engineered in order to alter a coding sequence for a variety of reasons, including but not limited to, alterations which modify the sequence to facilitate cloning, processing and/or expression of the gene product. For example, alterations are optionally introduced using techniques which are well known in the art, e.g., site-directed mutagenesis, to insert new restriction sites, to alter glycosylation patterns, to change codon preference, to introduce splice sites, etc.

Furthermore, a fragment or domain derived from any of the polypeptides of the invention can be combined with domains derived from other transcription factors or synthetic domains to modify the biological activity of a transcription factor. For instance, a DNA binding domain derived from a transcription factor of the invention can be combined with the activation domain of another transcription factor or with a synthetic activation domain. A transcription activation domain assists in initiating transcription from a DNA binding site. Examples include the transcription activation region of VP16 or GAL4 (Moore et al. (1998) Proc. Natl. Acad. Sci. USA 95: 376-381; and Aoyama et al. (1995) Plant Cell 7:1773-1785), peptides derived from bacterial sequences (Ma and Ptashne (1987) Cell 51: 113-119) and synthetic peptides (Giniger and Ptashne, (1987) Nature 330:670-672).

EXPRESSION AND MODIFICATION OF POLYPEPTIDES

Typically, polynucleotide sequences of the invention are incorporated into recombinant DNA (or RNA) molecules that direct expression of polypeptides of the invention in appropriate host cells, transgenic plants, in vitro translation systems, or the like. Due to the inherent degeneracy of the genetic code, nucleic acid sequences which encode substantially the same or a functionally equivalent amino acid sequence can be substituted for any listed sequence to provide for cloning and expressing the relevant homologue.

Vectors, Promoters and Expression Systems

The present invention includes recombinant constructs comprising one or more of the nucleic acid sequences herein. The constructs typically comprise a vector, such as a plasmid, a cosmid, a phage, a virus (e.g., a plant virus), a bacterial artificial chromosome (BAC), a yeast artificial chromosome (YAC), or the like, into which a nucleic acid sequence of the invention has been inserted, in a forward or reverse orientation. In a preferred aspect of this embodiment, the construct further comprises regulatory sequences, including, for example, a promoter, operably linked to the sequence. Large numbers of suitable vectors and promoters are known to those of skill in the art, and are commercially available.

General texts which describe molecular biological techniques useful herein, including the use and production of vectors, promoters and many other relevant topics, include Berger, Sambrook and Ausubel, *supra*. Any of the identified sequences can be incorporated into a cassette or vector, e.g., for expression in plants. A number of expression vectors suitable for stable transformation of plant cells or for the establishment of transgenic plants have been described including those described in Weissbach and Weissbach, (1989) Methods for Plant Molecular Biology, Academic Press, and Gelvin et al., (1990) Plant Molecular Biology Manual, Kluwer Academic Publishers. Specific examples include those derived from a Ti plasmid of *Agrobacterium tumefaciens*, as well as those disclosed by Herrera-Estrella et al. (1983) Nature 303: 209, Bevan (1984) Nucl Acid Res. 12: 8711-8721, Klee (1985) Bio/Technology 3: 637-642, for dicotyledonous plants.

Alternatively, non-Ti vectors can be used to transfer the DNA into monocotyledonous plants and cells by using free DNA delivery techniques. Such methods can involve, for example, the use of liposomes, electroporation, microprojectile bombardment, silicon carbide whiskers, and viruses. By using these methods transgenic plants such as wheat, rice (Christou (1991) Bio/Technology 9: 957-962) and corn (Gordon-Kamm (1990) Plant Cell 2: 603-618) can be produced. An immature embryo can also be a good target tissue for monocots for direct DNA delivery techniques by using the particle gun (Weeks et al. (1993) Plant Physiol 102: 1077-1084; Vasil (1993) Bio/Technology 10: 667-674; Wan and Lemeaux (1994) Plant Physiol 104: 37-48, and for *Agrobacterium*-mediated DNA transfer (Ishida et al. (1996) Nature Biotech 14: 745-750).

Typically, plant transformation vectors include one or more cloned plant coding sequence (genomic or cDNA) under the transcriptional control of 5' and 3' regulatory sequences and a dominant selectable marker. Such plant transformation vectors typically also contain a promoter (e.g., a regulatory region controlling inducible or constitutive, environmentally-or developmentally-regulated, or cell- or tissue-specific expression), a transcription initiation start site, an RNA processing signal (such as intron splice sites), a transcription termination site, and/or a polyadenylation signal.

Examples of constitutive plant promoters which can be useful for expressing the TF sequence include: the cauliflower mosaic virus (CaMV) 35S promoter, which confers constitutive, high-level expression in most plant tissues (*see*, e.g., Odel et al. (1985) Nature 313:810); the nopaline synthase promoter (An et al. (1988) Plant Physiol 88:547); and the octopine synthase promoter (Fromm et al. (1989) Plant Cell 1: 977).

A variety of plant gene promoters that regulate gene expression in response to environmental, hormonal, chemical, developmental signals, and in a tissue-active manner can be used for expression of a TF sequence in plants. Choice of a promoter is based largely on the phenotype of interest and is determined by such factors as tissue (e.g., seed, fruit, root, pollen, vascular tissue, flower, carpel, etc.), inducibility (e.g., in response to wounding, heat, cold, drought, light, pathogens, etc.), timing, developmental stage, and the like. Numerous known promoters have been characterized and can favorably be employed to promote expression of a polynucleotide of the invention in a transgenic plant or cell of interest. For example, tissue specific promoters include: seed-specific promoters (such as the napin, phaseolin or DC3 promoter described in US Pat. No. 5,773,697), fruit-specific promoters that are active during fruit ripening (such as the dru 1 promoter (US Pat. No. 5,783,393), or the 2A11 promoter (US Pat. No. 4,943,674) and the tomato polygalacturonase promoter (Bird et al. (1988) Plant Mol Biol 11:651), root-specific promoters, such as those disclosed in US Patent Nos. 5,618,988, 5,837,848 and 5,905,186, pollen-active promoters such as PTA29, PTA26 and PTA13 (US Pat. No. 5,792,929), promoters active in vascular tissue (Ringli and Keller (1998) Plant Mol Biol 37:977-988), flower-specific (Kaiser et al. (1995) Plant Mol Biol 28:231-243), pollen (Baerson et al. (1994) Plant Mol Biol 26:1947-1959), carpels (Ohl et al. (1990) Plant Cell 2:837-848), pollen and ovules (Baerson et al. (1993) Plant Mol Biol 22:255-267), auxin-inducible promoters (such as that described in van der Kop et al. (1999) Plant Mol Biol 39:979-990 or Baumann et al. (1999) Plant Cell 11:323-334), cytokinin-inducible promoter (Guevara-Garcia (1998) Plant Mol Biol 38:743-753), promoters responsive to gibberellin (Shi et al. (1998) Plant Mol Biol 38:1053-1060, Willmott et al. (1998) 38:817-825) and the like. Additional promoters are those that elicit expression in response to heat (Ainley et al. (1993) Plant Mol Biol 22: 13-23), light (e.g., the pea rbcS-3A promoter, Kuhlemeier et al. (1989) Plant Cell 1:471, and the maize rbcS promoter, Schaffner and Sheen (1991) Plant Cell 3: 997); wounding (e.g., *wun1*, Siebertz et al. (1989) Plant Cell 1: 961); pathogens (such as the PR-1 promoter described in Buchel et al. (1999) Plant Mol. Biol. 40:387-396, and the PDF1.2 promoter described in Manners et al. (1998) Plant Mol. Biol. 38:1071-80), and chemicals such as methyl jasmonate or salicylic acid (Gatz et al. (1997) Plant Mol Biol 48: 89-108). In addition, the timing of the expression can be controlled by using promoters such as those acting at senescence (An and Amazon (1995) Science 270: 1986-1988); or late seed development (Odell et al. (1994) Plant Physiol 106:447-458).

Plant expression vectors can also include RNA processing signals that can be positioned within, upstream or downstream of the coding sequence. In addition, the expression vectors can include additional regulatory sequences from the 3'-untranslated region of plant

genes, e.g., a 3' terminator region to increase mRNA stability of the mRNA, such as the PI-II terminator region of potato or the octopine or nopaline synthase 3' terminator regions.

Additional Expression Elements

Specific initiation signals can aid in efficient translation of coding sequences.

- 5 These signals can include, e.g., the ATG initiation codon and adjacent sequences. In cases where a coding sequence, its initiation codon and upstream sequences are inserted into the appropriate expression vector, no additional translational control signals may be needed. However, in cases where only coding sequence (e.g., a mature protein coding sequence), or a portion thereof, is inserted, exogenous transcriptional control signals including the ATG initiation codon can be
- 10 separately provided. The initiation codon is provided in the correct reading frame to facilitate transcription. Exogenous transcriptional elements and initiation codons can be of various origins, both natural and synthetic. The efficiency of expression can be enhanced by the inclusion of enhancers appropriate to the cell system in use.

Expression Hosts

- 15 The present invention also relates to host cells which are transduced with vectors of the invention, and the production of polypeptides of the invention (including fragments thereof) by recombinant techniques. Host cells are genetically engineered (i.e, nucleic acids are introduced, e.g., transduced, transformed or transfected) with the vectors of this invention, which may be, for example, a cloning vector or an expression vector comprising the relevant nucleic
- 20 acids herein. The vector is optionally a plasmid, a viral particle, a phage, a naked nucleic acids, *etc.* The engineered host cells can be cultured in conventional nutrient media modified as appropriate for activating promoters, selecting transformants, or amplifying the relevant gene. The culture conditions, such as temperature, pH and the like, are those previously used with the host cell selected for expression, and will be apparent to those skilled in the art and in the
- 25 references cited herein, including, Sambrook and Ausubel.

- The host cell can be a eukaryotic cell, such as a yeast cell, or a plant cell, or the host cell can be a prokaryotic cell, such as a bacterial cell. Plant protoplasts are also suitable for some applications. For example, the DNA fragments are introduced into plant tissues, cultured plant cells or plant protoplasts by standard methods including electroporation (Fromm et al.,
- 30 (1985) Proc. Natl. Acad. Sci. USA 82, 5824, infection by viral vectors such as cauliflower mosaic virus (CaMV) (Hohn et al., (1982) Molecular Biology of Plant Tumors, (Academic Press, New York) pp. 549-560; US 4,407,956), high velocity ballistic penetration by small particles with the nucleic acid either within the matrix of small beads or particles, or on the surface (Klein et al., (1987) Nature 327, 70-73), use of pollen as vector (WO 85/01856), or use of *Agrobacterium*

tumefaciens or *A. rhizogenes* carrying a T-DNA plasmid in which DNA fragments are cloned. The T-DNA plasmid is transmitted to plant cells upon infection by *Agrobacterium tumefaciens*, and a portion is stably integrated into the plant genome (Horsch et al. (1984) Science 233:496-498; Fraley et al. (1983) Proc. Natl. Acad. Sci. USA 80, 4803).

5 The cell can include a nucleic acid of the invention which encodes a polypeptide, wherein the cells expresses a polypeptide of the invention. The cell can also include vector sequences, or the like. Furthermore, cells and transgenic plants which include any polypeptide or nucleic acid above or throughout this specification, e.g., produced by transduction of a vector of the invention, are an additional feature of the invention.

10 For long-term, high-yield production of recombinant proteins, stable expression can be used. Host cells transformed with a nucleotide sequence encoding a polypeptide of the invention are optionally cultured under conditions suitable for the expression and recovery of the encoded protein from cell culture. The protein or fragment thereof produced by a recombinant cell may be secreted, membrane-bound, or contained intracellularly, depending on the sequence
15 and/or the vector used. As will be understood by those of skill in the art, expression vectors containing polynucleotides encoding mature proteins of the invention can be designed with signal sequences which direct secretion of the mature polypeptides through a prokaryotic or eukaryotic cell membrane.

Modified Amino Acids

20 Polypeptides of the invention may contain one or more modified amino acids. The presence of modified amino acids may be advantageous in, for example, increasing polypeptide half-life, reducing polypeptide antigenicity or toxicity, increasing polypeptide storage stability, or the like. Amino acid(s) are modified, for example, co-translationally or post-translationally during recombinant production or modified by synthetic or chemical means.

25 Non-limiting examples of a modified amino acid include incorporation or other use of acetylated amino acids, glycosylated amino acids, sulfated amino acids, prenylated (e.g., farnesylated, geranylgeranylated) amino acids, PEG modified (e.g., "PEGylated") amino acids, biotinylated amino acids, carboxylated amino acids, phosphorylated amino acids, etc. References adequate to guide one of skill in the modification of amino acids are replete throughout the
30 literature.

IDENTIFICATION OF ADDITIONAL FACTORS

A transcription factor provided by the present invention can also be used to identify additional endogenous or exogenous molecules that can affect a phenotype or trait of

interest. On the one hand, such molecules include organic (small or large molecules) and/or inorganic compounds that affect expression of (i.e., regulate) a particular transcription factor. Alternatively, such molecules include endogenous molecules that are acted upon either at a transcriptional level by a transcription factor of the invention to modify a phenotype as desired.

5 For example, the transcription factors can be employed to identify one or more downstream gene with which is subject to a regulatory effect of the transcription factor. In one approach, a transcription factor or transcription factor homologue of the invention is expressed in a host cell, e.g., a transgenic plant cell, tissue or explant, and expression products, either RNA or protein, of likely or random targets are monitored, e.g., by hybridization to a microarray of nucleic acid
10 probes corresponding to genes expressed in a tissue or cell type of interest, by two-dimensional gel electrophoresis of protein products, or by any other method known in the art for assessing expression of gene products at the level of RNA or protein. Alternatively, a transcription factor of the invention can be used to identify promoter sequences (i.e., binding sites) involved in the regulation of a downstream target. After identifying a promoter sequence, interactions between
15 the transcription factor and the promoter sequence can be modified by changing specific nucleotides in the promoter sequence or specific amino acids in the transcription factor that interact with the promoter sequence to alter a plant trait. Typically, transcription factor DNA binding sites are identified by gel shift assays. After identifying the promoter regions, the promoter region sequences can be employed in double-stranded DNA arrays to identify
20 molecules that affect the interactions of the transcription factors with their promoters (Bulyk et al. (1999) Nature Biotechnology 17:573-577).

The identified transcription factors are also useful to identify proteins that modify the activity of the transcription factor. Such modification can occur by covalent modification, such as by phosphorylation, or by protein-protein (homo or-heteropolymer) interactions. Any
25 method suitable for detecting protein-protein interactions can be employed. Among the methods that can be employed are co-immunoprecipitation, cross-linking and co-purification through gradients or chromatographic columns, and the two-hybrid yeast system.

The two-hybrid system detects protein interactions in vivo and is described in Chien, et al., (1991), Proc. Natl. Acad. Sci. USA 88, 9578-9582 and is commercially available
30 from Clontech (Palo Alto, Calif.). In such a system, plasmids are constructed that encode two hybrid proteins: one consists of the DNA-binding domain of a transcription activator protein fused to the TF polypeptide and the other consists of the transcription activator protein's activation domain fused to an unknown protein that is encoded by a cDNA that has been recombined into the plasmid as part of a cDNA library. The DNA-binding domain fusion plasmid

and the cDNA library are transformed into a strain of the yeast *Saccharomyces cerevisiae* that contains a reporter gene (e.g., lacZ) whose regulatory region contains the transcription activator's binding site. Either hybrid protein alone cannot activate transcription of the reporter gene.

Interaction of the two hybrid proteins reconstitutes the functional activator protein and results in expression of the reporter gene, which is detected by an assay for the reporter gene product. Then, the library plasmids responsible for reporter gene expression are isolated and sequenced to identify the proteins encoded by the library plasmids. After identifying proteins that interact with the transcription factors, assays for compounds that interfere with the TF protein-protein interactions can be preformed.

10 IDENTIFICATION OF MODULATORS

In addition to the intracellular molecules described above, extracellular molecules that alter activity or expression of a transcription factor, either directly or indirectly, can be identified. For example, the methods can entail first placing a candidate molecule in contact with a plant or plant cell. The molecule can be introduced by topical administration, such as spraying or soaking of a plant, and then the molecule's effect on the expression or activity of the TF polypeptide or the expression of the polynucleotide monitored. Changes in the expression of the TF polypeptide can be monitored by use of polyclonal or monoclonal antibodies, gel electrophoresis or the like. Changes in the expression of the corresponding polynucleotide sequence can be detected by use of microarrays, Northern, quantitative PCR, or any other technique for monitoring changes in mRNA expression. These techniques are exemplified in Ausubel et al. (eds) Current Protocols in Molecular Biology, John Wiley & Sons (1998). Such changes in the expression levels can be correlated with modified plant traits and thus identified molecules can be useful for soaking or spraying on fruit, vegetable and grain crops to modify traits in plants.

Essentially any available composition can be tested for modulatory activity of expression or activity of any nucleic acid or polypeptide herein. Thus, available libraries of compounds such as chemicals, polypeptides, nucleic acids and the like can be tested for modulatory activity. Often, potential modulator compounds can be dissolved in aqueous or organic (e.g., DMSO-based) solutions for easy delivery to the cell or plant of interest in which the activity of the modulator is to be tested. Optionally, the assays are designed to screen large modulator composition libraries by automating the assay steps and providing compounds from any convenient source to assays, which are typically run in parallel (e.g., in microtiter formats on microtiter plates in robotic assays).

In one embodiment, high throughput screening methods involve providing a combinatorial library containing a large number of potential compounds (potential modulator compounds). Such "combinatorial chemical libraries" are then screened in one or more assays, as described herein, to identify those library members (particular chemical species or subclasses) that display a desired characteristic activity. The compounds thus identified can serve as target compounds.

A combinatorial chemical library can be, e.g., a collection of diverse chemical compounds generated by chemical synthesis or biological synthesis. For example, a combinatorial chemical library such as a polypeptide library is formed by combining a set of chemical building blocks (e.g., in one example, amino acids) in every possible way for a given compound length (i.e., the number of amino acids in a polypeptide compound of a set length). Exemplary libraries include peptide libraries, nucleic acid libraries, antibody libraries (see, e.g., Vaughn et al. (1996) Nature Biotechnology, 14(3):309-314 and PCT/US96/10287), carbohydrate libraries (see, e.g., Liang et al. Science (1996) 274:1520-1522 and U.S. Patent 5,593,853), peptide nucleic acid libraries (see, e.g., U.S. Patent 5,539,083), and small organic molecule libraries (see, e.g., benzodiazepines, Baum C&EN Jan 18, page 33 (1993); isoprenoids, U.S. Patent 5,569,588; thiazolidinones and metathiazanones, U.S. Patent 5,549,974; pyrrolidines, U.S. Patents 5,525,735 and 5,519,134; morpholino compounds, U.S. Patent 5,506,337) and the like.

Preparation and screening of combinatorial or other libraries is well known to those of skill in the art. Such combinatorial chemical libraries include, but are not limited to, peptide libraries (see, e.g., U.S. Patent 5,010,175, Furka, Int. J. Pept. Prot. Res. 37:487-493 (1991) and Houghton et al. Nature 354:84-88 (1991)). Other chemistries for generating chemical diversity libraries can also be used.

In addition, as noted, compound screening equipment for high-throughput screening is generally available, e.g., using any of a number of well known robotic systems that have also been developed for solution phase chemistries useful in assay systems. These systems include automated workstations including an automated synthesis apparatus and robotic systems utilizing robotic arms. Any of the above devices are suitable for use with the present invention, e.g., for high-throughput screening of potential modulators. The nature and implementation of modifications to these devices (if any) so that they can operate as discussed herein will be apparent to persons skilled in the relevant art.

Indeed, entire high throughput screening systems are commercially available. These systems typically automate entire procedures including all sample and reagent pipetting, liquid dispensing, timed incubations, and final readings of the microplate in detector(s)

appropriate for the assay. These configurable systems provide high throughput and rapid start up as well as a high degree of flexibility and customization. Similarly, microfluidic implementations of screening are also commercially available.

The manufacturers of such systems provide detailed protocols the various high
5 throughput. Thus, for example, Zymark Corp. provides technical bulletins describing screening systems for detecting the modulation of gene transcription, ligand binding, and the like. The integrated systems herein, in addition to providing for sequence alignment and, optionally, synthesis of relevant nucleic acids, can include such screening apparatus to identify modulators that have an effect on one or more polynucleotides or polypeptides according to the present
10 invention.

In some assays it is desirable to have positive controls to ensure that the components of the assays are working properly. At least two types of positive controls are appropriate. That is, known transcriptional activators or inhibitors can be incubated with cells/plants/ etc. in one sample of the assay, and the resulting increase/decrease in transcription
15 can be detected by measuring the resulting increase in RNA/ protein expression, etc., according to the methods herein. It will be appreciated that modulators can also be combined with transcriptional activators or inhibitors to find modulators which inhibit transcriptional activation or transcriptional repression. Either expression of the nucleic acids and proteins herein or any additional nucleic acids or proteins activated by the nucleic acids or proteins herein, or both, can
20 be monitored.

In an embodiment, the invention provides a method for identifying compositions that modulate the activity or expression of a polynucleotide or polypeptide of the invention. For example, a test compound, whether a small or large molecule, is placed in contact with a cell, plant (or plant tissue or explant), or composition comprising the polynucleotide or polypeptide of
25 interest and a resulting effect on the cell, plant, (or tissue or explant) or composition is evaluated by monitoring, either directly or indirectly, one or more of: expression level of the polynucleotide or polypeptide, activity (or modulation of the activity) of the polynucleotide or polypeptide. In some cases, an alteration in a plant phenotype can be detected following contact of a plant (or plant cell, or tissue or explant) with the putative modulator, e.g., by modulation of expression or
30 activity of a polynucleotide or polypeptide of the invention.

SUBSEQUENCES

Also contemplated are uses of polynucleotides, also referred to herein as oligonucleotides, typically having at least 12 bases, preferably at least 15, more preferably at least

20, 30, or 50 bases, which hybridize under at least highly stringent (or ultra-high stringent or ultra-ultra- high stringent conditions) conditions to a polynucleotide sequence described above. The polynucleotides may be used as probes, primers, sense and antisense agents, and the like, according to methods as noted *supra*.

5 Subsequences of the polynucleotides of the invention, including polynucleotide fragments and oligonucleotides are useful as nucleic acid probes and primers. An oligonucleotide suitable for use as a probe or primer is at least about 15 nucleotides in length, more often at least about 18 nucleotides, often at least about 21 nucleotides, frequently at least about 30 nucleotides, or about 40 nucleotides, or more in length. A nucleic acid probe is useful in hybridization
10 protocols, e.g., to identify additional polypeptide homologues of the invention, including protocols for microarray experiments. Primers can be annealed to a complementary target DNA strand by nucleic acid hybridization to form a hybrid between the primer and the target DNA strand, and then extended along the target DNA strand by a DNA polymerase enzyme. Primer pairs can be used for amplification of a nucleic acid sequence, e.g., by the polymerase chain
15 reaction (PCR) or other nucleic-acid amplification methods. See Sambrook and Ausubel, *supra*.

In addition, the invention includes an isolated or recombinant polypeptide including a subsequence of at least about 15 contiguous amino acids encoded by the recombinant or isolated polynucleotides of the invention. For example, such polypeptides, or domains or fragments thereof, can be used as immunogens, e.g., to produce antibodies specific for the
20 polypeptide sequence, or as probes for detecting a sequence of interest. A subsequence can range in size from about 15 amino acids in length up to and including the full length of the polypeptide.

PRODUCTION OF TRANSGENIC PLANTS

Modification of Traits

The polynucleotides of the invention are favorably employed to produce
25 transgenic plants with various traits, or characteristics, that have been modified in a desirable manner, e.g., to improve the seed characteristics of a plant. For example, alteration of expression levels or patterns (e.g., spatial or temporal expression patterns) of one or more of the transcription factors (or transcription factor homologues) of the invention, as compared with the levels of the same protein found in a wild type plant, can be used to modify a plant's traits. An illustrative
30 example of trait modification, improved sugar-sensing characteristics, by altering expression levels of a particular transcription factor is described further in the Examples and the Sequence Listing.

Antisense and Cosuppression Approaches

In addition to expression of the nucleic acids of the invention as gene replacement or plant phenotype modification nucleic acids, the nucleic acids are also useful for sense and anti-sense suppression of expression, e.g., to down-regulate expression of a nucleic acid of the invention, e.g., as a further mechanism for modulating plant phenotype. That is, the nucleic acids of the invention, or subsequences or anti-sense sequences thereof, can be used to block expression of naturally occurring homologous nucleic acids. A variety of sense and anti-sense technologies are known in the art, e.g., as set forth in Lichtenstein and Nellen (1997)

Antisense Technology: A Practical Approach IRL Press at Oxford University, Oxford, England.

In general, sense or anti-sense sequences are introduced into a cell, where they are optionally amplified, e.g., by transcription. Such sequences include both simple oligonucleotide sequences and catalytic sequences such as ribozymes.

For example, a reduction or elimination of expression (i.e., a “knock-out”) of a transcription factor or transcription factor homologue polypeptide in a transgenic plant, e.g., to modify a plant trait, can be obtained by introducing an antisense construct corresponding to the polypeptide of interest as a cDNA. For antisense suppression, the transcription factor or homologue cDNA is arranged in reverse orientation (with respect to the coding sequence) relative to the promoter sequence in the expression vector. The introduced sequence need not be the full length cDNA or gene, and need not be identical to the cDNA or gene found in the plant type to be transformed. Typically, the antisense sequence need only be capable of hybridizing to the target gene or RNA of interest. Thus, where the introduced sequence is of shorter length, a higher degree of homology to the endogenous transcription factor sequence will be needed for effective antisense suppression. While antisense sequences of various lengths can be utilized, preferably, the introduced antisense sequence in the vector will be at least 30 nucleotides in length, and improved antisense suppression will typically be observed as the length of the antisense sequence increases. Preferably, the length of the antisense sequence in the vector will be greater than 100 nucleotides. Transcription of an antisense construct as described results in the production of RNA molecules that are the reverse complement of mRNA molecules transcribed from the endogenous transcription factor gene in the plant cell.

Suppression of endogenous transcription factor gene expression can also be achieved using a ribozyme. Ribozymes are RNA molecules that possess highly specific endoribonuclease activity. The production and use of ribozymes are disclosed in U.S. Patent No. 4,987,071 and U.S. Patent No. 5,543,508. Synthetic ribozyme sequences including antisense RNAs can be used to confer RNA cleaving activity on the antisense RNA, such that endogenous

mRNA molecules that hybridize to the antisense RNA are cleaved, which in turn leads to an enhanced antisense inhibition of endogenous gene expression.

5 Vectors in which RNA encoded by a transcription factor or transcription factor
homologue cDNA is over-expressed can also be used to obtain co-suppression of a corresponding
endogenous gene, e.g., in the manner described in U.S. Patent No. 5,231,020 to Jorgensen. Such
co-suppression (also termed sense suppression) does not require that the entire transcription factor
cDNA be introduced into the plant cells, nor does it require that the introduced sequence be
exactly identical to the endogenous transcription factor gene of interest. However, as with
antisense suppression, the suppressive efficiency will be enhanced as specificity of hybridization
10 is increased, e.g., as the introduced sequence is lengthened, and/or as the sequence similarity
between the introduced sequence and the endogenous transcription factor gene is increased.

Vectors expressing an untranslatable form of the transcription factor mRNA, e.g.,
sequences comprising one or more stop codon, or nonsense mutation) can also be used to
suppress expression of an endogenous transcription factor, thereby reducing or eliminating it's
15 activity and modifying one or more traits. Methods for producing such constructs are described
in U.S. Patent No. 5,583,021. Preferably, such constructs are made by introducing a premature
stop codon into the transcription factor gene. Alternatively, a plant trait can be modified by gene
silencing using double-strand RNA (Sharp (1999) Genes and Development 13: 139-141).

Another method for abolishing the expression of a gene is by insertion
20 mutagenesis using the T-DNA of *Agrobacterium tumefaciens*. After generating the insertion
mutants, the mutants can be screened to identify those containing the insertion in a transcription
factor or transcription factor homologue gene. Plants containing a single transgene insertion
event at the desired gene can be crossed to generate homozygous plants for the mutation (Koncz
et al. (1992) Methods in Arabidopsis Research, World Scientific).

25 Alternatively, a plant phenotype can be altered by eliminating an endogenous
gene, such as a transcription factor or transcription factor homologue, e.g., by homologous
recombination (Kempin et al. (1997) Nature 389:802).

A plant trait can also be modified by using the cre-lox system (for example, as
described in US Pat. No. 5,658,772). A plant genome can be modified to include first and
30 second lox sites that are then contacted with a Cre recombinase. If the lox sites are in the same
orientation, the intervening DNA sequence between the two sites is excised. If the lox sites are in
the opposite orientation, the intervening sequence is inverted.

The polynucleotides and polypeptides of this invention can also be expressed in a
plant in the absence of an expression cassette by manipulating the activity or expression level of

the endogenous gene by other means. For example, by ectopically expressing a gene by T-DNA activation tagging (Ichikawa et al. (1997) Nature 390 698-701; Kakimoto et al. (1996) Science 274: 982-985). This method entails transforming a plant with a gene tag containing multiple transcriptional enhancers and once the tag has inserted into the genome, expression of a flanking gene coding sequence becomes deregulated. In another example, the transcriptional machinery in a plant can be modified so as to increase transcription levels of a polynucleotide of the invention (See, e.g., PCT Publications WO 96/06166 and WO 98/53057 which describe the modification of the DNA binding specificity of zinc finger proteins by changing particular amino acids in the DNA binding motif).

The transgenic plant can also include the machinery necessary for expressing or altering the activity of a polypeptide encoded by an endogenous gene, for example by altering the phosphorylation state of the polypeptide to maintain it in an activated state.

Transgenic plants (or plant cells, or plant explants, or plant tissues) incorporating the polynucleotides of the invention and/or expressing the polypeptides of the invention can be produced by a variety of well established techniques as described above. Following construction of a vector, most typically an expression cassette, including a polynucleotide, e.g., encoding a transcription factor or transcription factor homologue, of the invention, standard techniques can be used to introduce the polynucleotide into a plant, a plant cell, a plant explant or a plant tissue of interest. Optionally, the plant cell, explant or tissue can be regenerated to produce a transgenic plant.

The plant can be any higher plant, including gymnosperms, monocotyledonous and dicotyledonous plants. Suitable protocols are available for *Leguminosae* (alfalfa, soybean, clover, etc.), *Umbelliferae* (carrot, celery, parsnip), *Cruciferae* (cabbage, radish, rapeseed, broccoli, etc.), *Curcubitaceae* (melons and cucumber), *Gramineae* (wheat, corn, rice, barley, millet, etc.), *Solanaceae* (potato, tomato, tobacco, peppers, etc.), and various other crops. See protocols described in Ammirato et al. (1984) Handbook of Plant Cell Culture –Crop Species. Macmillan Publ. Co. Shimamoto et al. (1989) Nature 338:274-276; Fromm et al. (1990) Bio/Technology 8:833-839; and Vasil et al. (1990) Bio/Technology 8:429-434.

Transformation and regeneration of both monocotyledonous and dicotyledonous plant cells is now routine, and the selection of the most appropriate transformation technique will be determined by the practitioner. The choice of method will vary with the type of plant to be transformed; those skilled in the art will recognize the suitability of particular methods for given plant types. Suitable methods can include, but are not limited to: electroporation of plant protoplasts; liposome-mediated transformation; polyethylene glycol (PEG) mediated

transformation; transformation using viruses; micro-injection of plant cells; micro-projectile bombardment of plant cells; vacuum infiltration; and *Agrobacterium tumefaciens* mediated transformation. Transformation means introducing a nucleotide sequence in a plant in a manner to cause stable or transient expression of the sequence.

5 Successful examples of the modification of plant characteristics by transformation with cloned sequences which serve to illustrate the current knowledge in this field of technology, and which are herein incorporated by reference, include: U.S. Patent Nos. 5,571,706; 5,677,175; 5,510,471; 5,750,386; 5,597,945; 5,589,615; 5,750,871; 5,268,526; 5,780,708; 5,538,880; 5,773,269; 5,736,369 and 5,610,042.

10 Following transformation, plants are preferably selected using a dominant selectable marker incorporated into the transformation vector. Typically, such a marker will confer antibiotic or herbicide resistance on the transformed plants, and selection of transformants can be accomplished by exposing the plants to appropriate concentrations of the antibiotic or herbicide.

15 After transformed plants are selected and grown to maturity, those plants showing a modified trait are identified. The modified trait can be any of those traits described above. Additionally, to confirm that the modified trait is due to changes in expression levels or activity of the polypeptide or polynucleotide of the invention can be determined by analyzing mRNA expression using Northern blots, RT-PCR or microarrays, or protein expression using
20 immunoblots or Western blots or gel shift assays.

INTEGRATED SYSTEMS—SEQUENCE IDENTITY

 Additionally, the present invention may be an integrated system, computer or computer readable medium that comprises an instruction set for determining the identity of one or more sequences in a database. In addition, the instruction set can be used to generate or identify
25 sequences that meet any specified criteria. Furthermore, the instruction set may be used to associate or link certain functional benefits, such improved sugar-sensing characteristics, with one or more identified sequence.

 For example, the instruction set can include, e.g., a sequence comparison or other alignment program, e.g., an available program such as, for example, the Wisconsin Package
30 Version 10.0, such as BLAST, FASTA, PILEUP, FINDPATTERNS or the like (GCG, Madison, WI). Public sequence databases such as GenBank, EMBL, Swiss-Prot and PIR or private sequence databases such as PhytoSeq (Incyte Pharmaceuticals, Palo Alto, CA) can be searched.

Alignment of sequences for comparison can be conducted by the local homology algorithm of Smith and Waterman (1981) Adv. Appl. Math. 2:482, by the homology alignment algorithm of Needleman and Wunsch (1970) J. Mol. Biol. 48:443, by the search for similarity method of Pearson and Lipman (1988) Proc. Natl. Acad. Sci. U.S.A. 85: 2444, by computerized
5 implementations of these algorithms. After alignment, sequence comparisons between two (or more) polynucleotides or polypeptides are typically performed by comparing sequences of the two sequences over a comparison window to identify and compare local regions of sequence similarity. The comparison window can be a segment of at least about 20 contiguous positions, usually about 50 to about 200, more usually about 100 to about 150 contiguous positions. A
10 description of the method is provided in Ausubel et al., *supra*.

A variety of methods of determining sequence relationships can be used, including manual alignment and computer assisted sequence alignment and analysis. This later approach is a preferred approach in the present invention, due to the increased throughput afforded by computer assisted methods. As noted above, a variety of computer programs for
15 performing sequence alignment are available, or can be produced by one of skill.

One example algorithm that is suitable for determining percent sequence identity and sequence similarity is the BLAST algorithm, which is described in Altschul et al. J. Mol. Biol 215:403-410 (1990). Software for performing BLAST analyses is publicly available, e.g., through the National Center for Biotechnology Information (<http://www.ncbi.nlm.nih.gov/>). This
20 algorithm involves first identifying high scoring sequence pairs (HSPs) by identifying short words of length W in the query sequence, which either match or satisfy some positive-valued threshold score T when aligned with a word of the same length in a database sequence. T is referred to as the neighborhood word score threshold (Altschul et al., *supra*). These initial neighborhood word hits act as seeds for initiating searches to find longer HSPs containing them.
25 The word hits are then extended in both directions along each sequence for as far as the cumulative alignment score can be increased. Cumulative scores are calculated using, for nucleotide sequences, the parameters M (reward score for a pair of matching residues; always > 0) and N (penalty score for mismatching residues; always < 0). For amino acid sequences, a scoring matrix is used to calculate the cumulative score. Extension of the word hits in each
30 direction are halted when: the cumulative alignment score falls off by the quantity X from its maximum achieved value; the cumulative score goes to zero or below, due to the accumulation of one or more negative-scoring residue alignments; or the end of either sequence is reached. The BLAST algorithm parameters W, T, and X determine the sensitivity and speed of the alignment. The BLASTN program (for nucleotide sequences) uses as defaults a wordlength (W) of 11, an

expectation (E) of 10, a cutoff of 100, M=5, N=-4, and a comparison of both strands. For amino acid sequences, the BLASTP program uses as defaults a wordlength (W) of 3, an expectation (E) of 10, and the BLOSUM62 scoring matrix (*see* Henikoff & Henikoff (1989) Proc. Natl. Acad. Sci. USA 89:10915).

5 In addition to calculating percent sequence identity, the BLAST algorithm also performs a statistical analysis of the similarity between two sequences (*see*, e.g., Karlin & Altschul (1993) Proc. Natl. Acad. Sci. USA 90:5873-5787). One measure of similarity provided by the BLAST algorithm is the smallest sum probability (P(N)), which provides an indication of the probability by which a match between two nucleotide or amino acid sequences would occur
10 by chance. For example, a nucleic acid is considered similar to a reference sequence (and, therefore, in this context, homologous) if the smallest sum probability in a comparison of the test nucleic acid to the reference nucleic acid is less than about 0.1, or less than about 0.01, and or even less than about 0.001. An additional example of a useful sequence alignment algorithm is PILEUP. PILEUP creates a multiple sequence alignment from a group of related sequences using
15 progressive, pairwise alignments. The program can align, e.g., up to 300 sequences of a maximum length of 5,000 letters.

 The integrated system, or computer typically includes a user input interface allowing a user to selectively view one or more sequence records corresponding to the one or more character strings, as well as an instruction set which aligns the one or more character strings
20 with each other or with an additional character string to identify one or more region of sequence similarity. The system may include a link of one or more character strings with a particular phenotype or gene function. Typically, the system includes a user readable output element which displays an alignment produced by the alignment instruction set.

 The methods of this invention can be implemented in a localized or distributed
25 computing environment. In a distributed environment, the methods may implemented on a single computer comprising multiple processors or on a multiplicity of computers. The computers can be linked, e.g. through a common bus, but more preferably the computer(s) are nodes on a network. The network can be a generalized or a dedicated local or wide-area network and, in certain preferred embodiments, the computers may be components of an intra-net or an internet.

30 Thus, the invention provides methods for identifying a sequence similar or homologous to one or more polynucleotides as noted herein, or one or more target polypeptides encoded by the polynucleotides, or otherwise noted herein and may include linking or associating a given plant phenotype or gene function with a sequence. In the methods, a sequence database is

provided (locally or across an inter or intra net) and a query is made against the sequence database using the relevant sequences herein and associated plant phenotypes or gene functions.

Any sequence herein can be entered into the database, before or after querying the database. This provides for both expansion of the database and, if done before the querying step, for insertion of control sequences into the database. The control sequences can be detected by the query to ensure the general integrity of both the database and the query. As noted, the query can be performed using a web browser based interface. For example, the database can be a centralized public database such as those noted herein, and the querying can be done from a remote terminal or computer across an internet or intranet.

EXAMPLES

The following examples are intended to illustrate but not limit the present invention.

EXAMPLE I. FULL LENGTH GENE IDENTIFICATION AND CLONING

Putative transcription factor sequences (genomic or ESTs) related to known transcription factors were identified in the *Arabidopsis thaliana* GenBank database using the tblastn sequence analysis program using default parameters and a P-value cutoff threshold of -4 or -5 or lower, depending on the length of the query sequence. Putative transcription factor sequence hits were then screened to identify those containing particular sequence strings. If the sequence hits contained such sequence strings, the sequences were confirmed as transcription factors.

Alternatively, *Arabidopsis thaliana* cDNA libraries derived from different tissues or treatments, or genomic libraries were screened to identify novel members of a transcription family using a low stringency hybridization approach. Probes were synthesized using gene specific primers in a standard PCR reaction (annealing temperature 60° C) and labeled with ³²P dCTP using the High Prime DNA Labeling Kit (Boehringer Mannheim). Purified radiolabelled probes were added to filters immersed in Church hybridization medium (0.5 M NaPO₄ pH 7.0, 7% SDS, 1 % w/v bovine serum albumin) and hybridized overnight at 60 °C with shaking. Filters were washed two times for 45 to 60 minutes with 1xSCC, 1% SDS at 60° C.

To identify additional sequence 5' or 3' of a partial cDNA sequence in a cDNA library, 5' and 3' rapid amplification of cDNA ends (RACE) was performed using the Marathon™ cDNA amplification kit (Clontech, Palo Alto, CA). Generally, the method entailed first isolating poly(A) mRNA, performing first and second strand cDNA synthesis to generate double stranded

cDNA, blunting cDNA ends, followed by ligation of the Marathon™ Adaptor to the cDNA to form a library of adaptor-ligated ds cDNA.

Gene-specific primers were designed to be used along with adaptor specific primers for both 5' and 3' RACE reactions. Nested primers, rather than single primers, were used to increase PCR specificity. Using 5' and 3' RACE reactions, 5' and 3' RACE fragments were obtained, sequenced and cloned. The process can be repeated until 5' and 3' ends of the full-length gene were identified. Then the full-length cDNA was generated by PCR using primers specific to 5' and 3' ends of the gene by end-to-end PCR.

EXAMPLE II. CONSTRUCTION OF EXPRESSION VECTORS

The sequence was amplified from a genomic or cDNA library using primers specific to sequences upstream and downstream of the coding region. The expression vector was pMEN20 or pMEN65, which are both derived from pMON316 (Sanders et al, (1987) Nucleic Acids Research 15:1543-58) and contain the CaMV 35S promoter to express transgenes. To clone the sequence into the vector, both pMEN20 and the amplified DNA fragment were digested separately with SalI and NotI restriction enzymes at 37° C for 2 hours. The digestion products were subject to electrophoresis in a 0.8% agarose gel and visualized by ethidium bromide staining. The DNA fragments containing the sequence and the linearized plasmid were excised and purified by using a Qiaquick gel extraction kit (Qiagen, CA). The fragments of interest were ligated at a ratio of 3:1 (vector to insert). Ligation reactions using T4 DNA ligase (New England Biolabs, MA) were carried out at 16° C for 16 hours. The ligated DNAs were transformed into competent cells of the *E. coli* strain DH5alpha by using the heat shock method. The transformations were plated on LB plates containing 50 mg/l kanamycin (Sigma).

Individual colonies were grown overnight in five milliliters of LB broth containing 50 mg/l kanamycin at 37° C. Plasmid DNA was purified by using Qiaquick Mini Prep kits (Qiagen, CA).

EXAMPLE III. TRANSFORMATION OF AGROBACTERIUM WITH THE EXPRESSION VECTOR

After the plasmid vector containing the gene was constructed, the vector was used to transform *Agrobacterium tumefaciens* cells expressing the gene products. The stock of *Agrobacterium tumefaciens* cells for transformation were made as described by Nagel et al. (1990) FEMS Microbiol Letts. 67: 325-328. *Agrobacterium* strain ABI was grown in 250 ml LB medium (Sigma) overnight at 28°C with shaking until an absorbance (A_{600}) of 0.5 – 1.0 was reached. Cells were harvested by centrifugation at 4,000 x g for 15 min at 4° C. Cells were then

resuspended in 250 µl chilled buffer (1 mM HEPES, pH adjusted to 7.0 with KOH). Cells were centrifuged again as described above and resuspended in 125 µl chilled buffer. Cells were then centrifuged and resuspended two more times in the same HEPES buffer as described above at a volume of 100 µl and 750 µl, respectively. Resuspended cells were then distributed into 40 µl aliquots, quickly frozen in liquid nitrogen, and stored at -80° C.

Agrobacterium cells were transformed with plasmids prepared as described above following the protocol described by Nagel et al. For each DNA construct to be transformed, 50 – 100 ng DNA (generally resuspended in 10 mM Tris-HCl, 1 mM EDTA, pH 8.0) was mixed with 40 µl of *Agrobacterium* cells. The DNA/cell mixture was then transferred to a chilled cuvette with a 2mm electrode gap and subject to a 2.5 kV charge dissipated at 25 µF and 200 µF using a Gene Pulser II apparatus (Bio-Rad). After electroporation, cells were immediately resuspended in 1.0 ml LB and allowed to recover without antibiotic selection for 2 – 4 hours at 28° C in a shaking incubator. After recovery, cells were plated onto selective medium of LB broth containing 100 µg/ml spectinomycin (Sigma) and incubated for 24-48 hours at 28° C. Single colonies were then picked and inoculated in fresh medium. The presence of the plasmid construct was verified by PCR amplification and sequence analysis.

EXAMPLE IV. TRANSFORMATION OF *ARABIDOPSIS* PLANTS WITH *AGROBACTERIUM TUMEFACIENS* WITH EXPRESSION VECTOR

After transformation of *Agrobacterium tumefaciens* with plasmid vectors containing the gene, single *Agrobacterium* colonies were identified, propagated, and used to transform *Arabidopsis* plants. Briefly, 500 ml cultures of LB medium containing 50 mg/l kanamycin were inoculated with the colonies and grown at 28° C with shaking for 2 days until an absorbance (A_{600}) of > 2.0 is reached. Cells were then harvested by centrifugation at 4,000 x g for 10 min, and resuspended in infiltration medium (1/2 X Murashige and Skoog salts (Sigma), 1 X Gamborg's B-5 vitamins (Sigma), 5.0% (w/v) sucrose (Sigma), 0.044 µM benzylamino purine (Sigma), 200 µl/L Silwet L-77 (Lehle Seeds) until an absorbance (A_{600}) of 0.8 was reached.

Prior to transformation, *Arabidopsis thaliana* seeds (ecotype Columbia) were sown at a density of ~10 plants per 4" pot onto Pro-Mix BX potting medium (Hummert International) covered with fiberglass mesh (18 mm X 16 mm). Plants were grown under continuous illumination (50-75 µE/m²/sec) at 22-23° C with 65-70% relative humidity. After about 4 weeks, primary inflorescence stems (bolts) are cut off to encourage growth of multiple secondary bolts. After flowering of the mature secondary bolts, plants were prepared for transformation by removal of all siliques and opened flowers.

The pots were then immersed upside down in the mixture of *Agrobacterium* infiltration medium as described above for 30 sec, and placed on their sides to allow draining into a 1' x 2' flat surface covered with plastic wrap. After 24 h, the plastic wrap was removed and pots are turned upright. The immersion procedure was repeated one week later, for a total of two
5 immersions per pot. Seeds were then collected from each transformation pot and analyzed following the protocol described below.

EXAMPLE V. IDENTIFICATION OF ARABIDOPSIS PRIMARY TRANSFORMANTS

Seeds collected from the transformation pots were sterilized essentially as follows. Seeds were dispersed into in a solution containing 0.1% (v/v) Triton X-100 (Sigma) and
10 sterile H₂O and washed by shaking the suspension for 20 min. The wash solution was then drained and replaced with fresh wash solution to wash the seeds for 20 min with shaking. After removal of the second wash solution, a solution containing 0.1% (v/v) Triton X-100 and 70% ethanol (Equistar) was added to the seeds and the suspension was shaken for 5 min. After
15 removal of the ethanol/detergent solution, a solution containing 0.1% (v/v) Triton X-100 and 30% (v/v) bleach (Clorox) was added to the seeds, and the suspension was shaken for 10 min. After removal of the bleach/detergent solution, seeds were then washed five times in sterile distilled H₂O. The seeds were stored in the last wash water at 4°C for 2 days in the dark before being
20 plated onto antibiotic selection medium (1 X Murashige and Skoog salts (pH adjusted to 5.7 with 1M KOH), 1 X Gamborg's B-5 vitamins, 0.9% phytagar (Life Technologies), and 50 mg/l kanamycin). Seeds were germinated under continuous illumination (50-75 µE/m²/sec) at 22-23°C. After 7-10 days of growth under these conditions, kanamycin resistant primary transformants (T₁ generation) were visible and obtained. These seedlings were transferred first to fresh
selection plates where the seedlings continued to grow for 3-5 more days, and then to soil (Pro-Mix BX potting medium).

25 Primary transformants were crossed and progeny seeds (T₂) collected; kanamycin resistant seedlings were selected and analyzed. The expression levels of the recombinant polynucleotides in the transformants varies from about a 5% expression level increase to a least a 100% expression level increase. Similar observations are made with respect to polypeptide level expression.

30

EXAMPLE VI. IDENTIFICATION OF ARABIDOPSIS PLANTS WITH TRANSCRIPTION FACTOR GENE KNOCKOUTS

The screening of insertion mutagenized *Arabidopsis* collections for null mutants in a known target gene was essentially as described in Krysan et al (1999) Plant Cell 11:2283-2290. Briefly, gene-specific primers, nested by 5-250 base pairs to each other, were designed from the 5' and 3' regions of a known target gene. Similarly, nested sets of primers were also created specific to each of the T-DNA or transposon ends (the "right" and "left" borders). All possible combinations of gene specific and T-DNA/transposon primers were used to detect by PCR an insertion event within or close to the target gene. The amplified DNA fragments were then sequenced which allows the precise determination of the T-DNA/transposon insertion point relative to the target gene. Insertion events within the coding or intervening sequence of the genes were deconvoluted from a pool comprising a plurality of insertion events to a single unique mutant plant for functional characterization. The method is described in more detail in Yu and Adam, US Application Serial No. 09/177,733 filed October 23, 1998.

EXAMPLE VII. IDENTIFICATION OF SUGAR-SENSING CHARACTERISTICS PHENOTYPE IN OVEREXPRESSOR OR GENE KNOCKOUT PLANTS

Experiments were performed to identify those transformants or knockouts that exhibited modified sugar-sensing. For such studies, seeds from transformants were germinated on media containing 5% glucose or 9.4% sucrose which normally partially restrict hypocotyl elongation. Plants with altered sugar sensing may have either longer or shorter hypocotyls than normal plants when grown on this media. Additionally, other plant traits may be varied such as root mass.

Table 3 shows the phenotypes observed for particular overexpressor or knockout plants and provides the SEQ ID No., the internal reference code (GID), whether a knockout or overexpressor plant was analyzed and the observed phenotype.

Table 3

SEQ ID No.	GID	Knockout (OE) or overexpressor KO)	Phenotype observed
1	G26	OE	Decreased germination and growth on glucose medium
3	G38	OE	Reduced germination on glucose medium
5	G43	OE	Decreased germination and growth on glucose medium
7	G207	OE	Decreased germination on glucose medium
9	G241	OE	Decreased germination and growth on glucose medium
11	G254	OE	Decreased germination and growth on glucose medium
13	G263	OE	Decreased root growth on sucrose medium
15	G308	OE	No germination on glucose medium
17	G536	OE	Decreased germination and growth on glucose medium
19	G680	OE	Reduced germination on glucose medium
21	G867	OE	Better seedling vigor on sucrose medium
23	G912	OE	Reduced cotyledon expansion in glucose
25	G996	OE	Reduced germination on glucose medium
27	G1068	OE	Reduced cotyledon expansion in glucose
29	G1337	OE	Decreased germination on sucrose medium

For a particular overexpressor that shows a less beneficial sugar-sensing characteristic, it may be more useful to select a plant with a decreased expression of the particular transcription factor. For a particular knockout that shows a less beneficial sugar-sensing characteristic, it may be more useful to select a plant with an increased expression of the particular transcription factor.

EXAMPLE VIII. IDENTIFICATION OF HOMOLOGOUS SEQUENCES

Homologous sequences from *Arabidopsis* and plant species other than *Arabidopsis* were identified using database sequence search tools, such as the Basic Local Alignment Search Tool (BLAST) (Altschul et al. (1990) *J. Mol. Biol.* 215:403-410; and Altschul et al. (1997) *Nucl. Acid Res.* 25: 3389-3402). The tblastx sequence analysis programs were employed using the BLOSUM-62 scoring matrix (Henikoff, S. and Henikoff, J. G. (1992) *Proc. Natl. Acad. Sci. USA* 89: 10915-10919).

Identified *Arabidopsis* homologous sequences are provided in Figure 2 and included in the Sequence Listing. The percent sequence identity among these sequences is as low as 47% sequence identity. Additionally, the entire NCBI GenBank database was filtered for sequences from all plants except *Arabidopsis thaliana* by selecting all entries in the NCBI GenBank database associated with NCBI taxonomic ID 33090 (Viridiplantae; all plants) and excluding entries associated with taxonomic ID 3701 (*Arabidopsis thaliana*). These sequences were compared to sequences representing genes of SEQ IDs Nos. 1-54 on 9/26/2000 using the Washington University TBLASTX algorithm (version 2.0a19MP). For each gene of SEQ IDs

Nos. 1-54, individual comparisons were ordered by probability score (P-value), where the score reflects the probability that a particular alignment occurred by chance. For example, a score of $3.6e-40$ is 3.6×10^{-40} . For up to ten species, the gene with the lowest P-value (and therefore the most likely homolog) is listed in Figure 3.

5 In addition to P-values, comparisons were also scored by percentage identity. Percentage identity reflects the degree to which two segments of DNA or protein are identical over a particular length. The ranges of percent identity between the non-Arabidopsis genes shown in Figure 3 and the Arabidopsis genes in the sequence listing are: SEQ ID No. 1: 44%-79%; SEQ ID No. 3: 36%-72%; SEQ ID No. 5: 42%-67%; SEQ ID No. 7: 55%-82%; SEQ ID No. 9: 69%-84%;
 10 SEQ ID No. 11: 57%-90%; SEQ ID No. 13: 48%-85%; SEQ ID No. 15: 38%-85%; SEQ ID No. 17: 77%-87%; SEQ ID No. 19: 42%-88%; SEQ ID No. 21: 54%-69%; SEQ ID No. 23: 34%-71%; SEQ ID No. 25: 55%-95%; SEQ ID No. 27: 54%-95%; SEQ ID No. 29: 37%-58%; SEQ ID No. 31: 42%-70%; SEQ ID No. 33: 46%-62%; SEQ ID No. 35: 64%-84%; SEQ ID No. 37: 57%-87%; SEQ ID No. 39: 40%-80%; SEQ ID No. 41: 56%-82%; SEQ ID No. 43: 64%-93%; SEQ ID
 15 No. 45: 35%-86%; SEQ ID No. 47: 84%-91%; SEQ ID No. 49: 85%-91%; SEQ ID No. 51: 38%-89%; SEQ ID No. 53: 53%-75%; SEQ ID No. 55: 57%-72%; SEQ ID No. 57: 57%-69%; SEQ ID No. 59: 49%-86%; SEQ ID No. 61: 49%-78%; SEQ ID No. 63: 51%-86%; SEQ ID No. 65: 42%-72%; SEQ ID No. 67: 35%-69%; and SEQ ID No. 69: 36%-64%.

20 The polynucleotides and polypeptides in the Sequence Listing and the identified homologous sequences may be stored in a computer system and have associated or linked with the sequences a function, such as that the polynucleotides and polypeptides are useful for modifying the sugar-sensing characteristics of a plant.

25 All references, publications, patents and other documents herein are incorporated by reference in their entirety for all purposes. Although the invention has been described with reference to the embodiments and examples above, it should be understood that various modifications can be made without departing from the spirit of the invention.

What is claimed is:

1. A transgenic plant with modified sugar-sensing characteristics, which plant comprises a recombinant polynucleotide comprising a nucleotide sequence selected from the group consisting of:

- 5 (a) a nucleotide sequence encoding a polypeptide comprising a sequence selected from SEQ ID Nos. 2N, where N=1-35, or a complementary nucleotide sequence thereof;
- (b) a nucleotide sequence encoding a polypeptide comprising a conservatively substituted variant of a polypeptide of (a);
- (c) a nucleotide sequence comprising a sequence selected from those of SEQ ID Nos. 2N-10 1, where N=1-35, or a complementary nucleotide sequence thereof;
- (d) a nucleotide sequence comprising silent substitutions in a nucleotide sequence of (c);
- (e) a nucleotide sequence which hybridizes under stringent conditions to a nucleotide sequence of one or more of: (a), (b), (c), or (d);
- (f) a nucleotide sequence comprising at least 15 consecutive nucleotides of a sequence of 15 any of (a)-(e);
- (g) a nucleotide sequence comprising a subsequence or fragment of any of (a)-(f), which subsequence or fragment encodes a polypeptide that modifies a plant's sugar-sensing characteristics;
- (h) a nucleotide sequence having at least 34% sequence identity to a nucleotide sequence 20 of any of (a)-(g);
- (i) a nucleotide sequence having at least 60% identity sequence identity to a nucleotide sequence of any of (a)-(g);
- (j) a nucleotide sequence which encodes a polypeptide having at least 34% identity sequence identity to a polypeptide of SEQ ID Nos. 2N, where N=1-35;
- 25 (k) a nucleotide sequence which encodes a polypeptide having at least 60% identity sequence identity to a polypeptide of SEQ ID Nos. 2N, where N=1-35; and
- (l) a nucleotide sequence which encodes a polypeptide having at least 65% sequence identity to a conserved domain of a polypeptide of SEQ ID Nos. 2N, where N=1-35.

30 2. The transgenic plant of claim 1, further comprising a constitutive, inducible, or tissue-active promoter operably linked to said nucleotide sequence.

3. The transgenic plant of claim 1, wherein the plant is selected from the group consisting of: soybean, wheat, corn, potato, cotton, rice, oilseed rape, sunflower, alfalfa, sugarcane, turf,

banana, blackberry, blueberry, strawberry, raspberry, cantaloupe, carrot, cauliflower, coffee, cucumber, eggplant, grapes, honeydew, lettuce, mango, melon, onion, papaya, peas, peppers, pineapple, spinach, squash, sweet corn, tobacco, tomato, watermelon, rosaceous fruits, and vegetable brassicas.

5

4. An isolated or recombinant polynucleotide comprising a nucleotide sequence selected from the group consisting of:

- (a) a nucleotide sequence encoding a polypeptide comprising a sequence selected from SEQ ID Nos. 2N, where N=1-35, or a complementary nucleotide sequence thereof;
- 10 (b) a nucleotide sequence encoding a polypeptide comprising a conservatively substituted variant of a polypeptide of (a);
- (c) a nucleotide sequence comprising a sequence selected from those of SEQ ID Nos. 2N-1, where N=1-35, or a complementary nucleotide sequence thereof;
- (d) a nucleotide sequence comprising silent substitutions in a nucleotide sequence of (c);
- 15 (e) a nucleotide sequence which hybridizes under stringent conditions to a nucleotide sequence of one or more of: (a), (b), (c), or (d);
- (f) a nucleotide sequence comprising at least 15 consecutive nucleotides of a sequence of any of (a)-(e);
- (g) a nucleotide sequence comprising a subsequence or fragment of any of (a)-(f), which
- 20 subsequence or fragment encodes a polypeptide that modifies a plant's sugar-sensing characteristics;
- (h) a nucleotide sequence having at least 34% sequence identity to a nucleotide sequence of any of (a)-(g);
- (i) a nucleotide sequence having at least 60% identity sequence identity to a nucleotide
- 25 sequence of any of (a)-(g);
- (j) a nucleotide sequence which encodes a polypeptide having at least 34% identity sequence identity to a polypeptide of SEQ ID Nos. 2N, where N=1-35;
- (k) a nucleotide sequence which encodes a polypeptide having at least 60% identity sequence identity to a polypeptide of SEQ ID Nos. 2N, where N=1-35; and
- 30 (l) a nucleotide sequence which encodes a conserved domain of a polypeptide having at least 65% sequence identity to a conserved domain of a polypeptide of SEQ ID Nos. 2N, where N=1-35.

5. The isolated or recombinant polynucleotide of claim 4, further comprising a constitutive, inducible, or tissue-active promoter operably linked to the nucleotide sequence.

6. A cloning or expression vector comprising the isolated or recombinant polynucleotide of claim 4.

7. A cell comprising the cloning or expression vector of claim 6.

8. A transgenic plant comprising the isolated or recombinant polynucleotide of claim 4.

10

9. A composition produced by one or more of:

(a) incubating one or more polynucleotide of claim 4 with a nuclease;

(b) incubating one or more polynucleotide of claim 4 with a restriction enzyme;

(c) incubating one or more polynucleotide of claim 4 with a polymerase;

15

(d) incubating one or more polynucleotide of claim 4 with a polymerase and a primer;

(e) incubating one or more polynucleotide of claim 4 with a cloning vector, or

(f) incubating one or more polynucleotide of claim 4 with a cell.

10. A composition comprising two or more different polynucleotides of claim 4.

20

11. An isolated or recombinant polypeptide comprising a subsequence of at least about 15 contiguous amino acids encoded by the recombinant or isolated polynucleotide of claim 4.

12. A plant ectopically expressing an isolated polypeptide of claim 11.

25

13. A method for producing a plant having modified sugar-sensing characteristics, the method comprising altering the expression of the isolated or recombinant polynucleotide of claim 4 or the expression levels or activity of a polypeptide of claim 11 in a plant, thereby producing a modified plant, and selecting the modified plant for improved sugar-sensing characteristics thereby providing the modified plant with a modified sugar-sensing characteristics.

30

14. The method of claim 13, wherein the polynucleotide is a polynucleotide of claim 4.

15. A method of identifying a factor that is modulated by or interacts with a polypeptide encoded by a polynucleotide of claim 4, the method comprising:

- (a) expressing a polypeptide encoded by the polynucleotide in a plant; and
- (b) identifying at least one factor that is modulated by or interacts with the polypeptide.

5

16. The method of claim 15, wherein the identifying is performed by detecting binding by the polypeptide to a promoter sequence, or detecting interactions between an additional protein and the polypeptide in a yeast two hybrid system.

10 17. The method of claim 15, wherein the identifying is performed by detecting expression of a factor by hybridization to a microarray, subtractive hybridization or differential display.

18. A method of identifying a molecule that modulates activity or expression of a polynucleotide or polypeptide of interest, the method comprising:

- 15 (a) placing the molecule in contact with a plant comprising the polynucleotide or polypeptide encoded by the polynucleotide of claim 4; and,
- (b) monitoring one or more of:
 - (i) expression level of the polynucleotide in the plant;
 - (ii) expression level of the polypeptide in the plant;
 - 20 (iii) modulation of an activity of the polypeptide in the plant; or
 - (iv) modulation of an activity of the polynucleotide in the plant.

19. An integrated system, computer or computer readable medium comprising one or more character strings corresponding to a polynucleotide of claim 4, or to a polypeptide encoded by the polynucleotide.

25

20. The integrated system, computer or computer readable medium of claim 19, further comprising a link between said one or more sequence strings to a modified plant sugar-sensing characteristics phenotype.

30

21. A method of identifying a sequence similar or homologous to one or more polynucleotides of claim 4, or one or more polypeptides encoded by the polynucleotides, the method comprising:

- (a) providing a sequence database; and,

(b) querying the sequence database with one or more target sequences corresponding to the one or more polynucleotides or to the one or more polypeptides to identify one or more sequence members of the database that display sequence similarity or homology to one or more of the one or more target sequences.

5

22. The method of claim 21, wherein the querying comprises aligning one or more of the target sequences with one or more of the one or more sequence members in the sequence database.

10

23. The method of claim 21, wherein the querying comprises identifying one or more of the one or more sequence members of the database that meet a user-selected identity criteria with one or more of the target sequences.

15

24. The method of claim 21, further comprising linking the one or more of the polynucleotides of claim 4, or encoded polypeptides, to a modified plant sugar-sensing characteristics phenotype.

20

25. A plant comprising altered expression levels of an isolated or recombinant polynucleotide of claim 4.

26. A plant comprising altered expression levels or the activity of an isolated or recombinant polypeptide of claim 11.

Figure 1

SEQ ID No.	GID	cDNA or protein	conserved domain
1	G26	cDNA	
2	G26	protein	67-134
3	G38	cDNA	
4	G38	protein	76-143
5	G43	cDNA	
6	G43	protein	104-172
7	G207	cDNA	
8	G207	protein	6-106
9	G241	cDNA	
10	G241	protein	14-114
11	G254	cDNA	
12	G254	protein	62-106
13	G263	cDNA	
14	G263	protein	15-105
15	G308	cDNA	
16	G308	protein	270-274
17	G536	cDNA	
18	G536	protein	226-233
19	G680	cDNA	
20	G680	protein	24-70
21	G867	cDNA	
22	G867	protein	59-124
23	G912	cDNA	
24	G912	protein	51-118
25	G996	cDNA	
26	G996	protein	14-114
27	G1068	cDNA	
28	G1068	protein	143-150
29	G1337	cDNA	
30	G1337	protein	9-75

Figure 2

SEQ ID No.	GID	homolog	cDNA or protein	conserved domain
31	G1141	homolog of G38	cDNA	
32	G1141	homolog of G38	protein	75-142
33	G46	homolog of G43	cDNA	
34	G46	homolog of G43	protein	107-175
35	G242	homolog of G207	cDNA	
36	G242	homolog of G207	protein	6-105
37	G227	homolog of G207	cDNA	
38	G227	homolog of G207	protein	13-112
39	G1307	homolog of G241	cDNA	
40	G1307	homolog of G241	protein	14-114
41	G1327	homolog of G241	cDNA	
42	G1327	homolog of G241	protein	14-116
43	G673	homolog of G254	cDNA	
44	G673	homolog of G254	protein	37-95
45	G307	homolog of G308	cDNA	
46	G307	homolog of G308	protein	323-339
47	G529	homolog of G536	cDNA	
48	G529	homolog of G536	protein	229-236
49	G531	homolog of G536	cDNA	
50	G531	homolog of G536	protein	227-234
51	G214	homolog of G680	cDNA	
52	G214	homolog of G680	protein	22-71
53	G1930	homolog of G867	cDNA	
54	G1930	homolog of G867	protein	59-124
55	G9	homolog of G867	cDNA	
56	G9	homolog of G867	protein	62-127
57	G993	homolog of G867	cDNA	
58	G993	homolog of G867	protein	69-134
59	G41	homolog of G912	cDNA	
60	G41	homolog of G912	protein	39-106
61	G40	homolog of G912	cDNA	
62	G40	homolog of G912	protein	45-112
63	G42	homolog of G912	cDNA	
64	G42	homolog of G912	protein	48-115
65	G1127	homolog of G1068	cDNA	
66	G1127	homolog of G1068	protein	103-110, 155-162
67	G2657	homolog of G1068	cDNA	
68	G2657	homolog of G1068	protein	116-129
69	G326	homolog of G1337	cDNA	
70	G326	homolog of G1337	protein	11-94, 354-400

Figure 3A

SEQ ID No.	GID	Genbank NID	P-value	Species
1	G26	4387560	6.00E-27	Lycopersicon esculentum
1	G26	9427282	1.60E-24	Triticum aestivum
1	G26	7206394	4.30E-24	Medicago truncatula
1	G26	7796858	5.40E-24	Glycine max
1	G26	7788764	6.40E-24	Lotus japonicus
1	G26	8098026	1.40E-20	Hordeum vulgare
1	G26	790362	1.80E-20	Nicotiana tabacum
1	G26	569065	2.60E-20	Oryza sativa
1	G26	3264766	3.60E-20	Prunus armeniaca
1	G26	7528275	6.20E-20	Mesembryanthemum crystallinum
3	G38	8346772	5.30E-47	Catharanthus roseus
3	G38	7205636	3.80E-45	Medicago truncatula
3	G38	7684799	2.80E-43	Glycine max
3	G38	9363798	4.80E-38	Triticum aestivum
3	G38	7777379	7.80E-38	Lotus japonicus
3	G38	8903111	1.10E-33	Hordeum vulgare
3	G38	568076	9.70E-28	Oryza sativa
3	G38	9434234	3.90E-19	Lycopersicon esculentum
3	G38	7324705	5.40E-16	Lycopersicon pennellii
3	G38	9298423	6.70E-16	Sorghum bicolor
5	G43	5760554	1.50E-29	Glycine max
5	G43	7778996	1.80E-29	Lotus japonicus
5	G43	5603736	5.00E-25	Lycopersicon esculentum
5	G43	6478844	1.00E-23	Matricaria chamomilla
5	G43	790361	1.60E-23	Nicotiana tabacum
5	G43	7528275	2.00E-23	Mesembryanthemum crystallinum
5	G43	9199136	2.30E-23	Medicago truncatula
5	G43	8980312	6.20E-23	Catharanthus roseus
5	G43	8809570	1.30E-22	Nicotiana sylvestris
5	G43	7627061	8.50E-22	Gossypium arboreum
7	G207	6529807	1.50E-63	Lycopersicon esculentum
7	G207	7564212	1.00E-57	Medicago truncatula
7	G207	7624453	1.60E-57	Gossypium arboreum
7	G207	5820271	6.50E-54	Glycine max
7	G207	7322467	3.40E-52	Lycopersicon hirsutum
7	G207	5045349	2.10E-46	Gossypium hirsutum
7	G207	8071527	2.60E-44	Solanum tuberosum
7	G207	7790004	3.70E-43	Beta vulgaris
7	G207	6325768	2.10E-40	Lotus japonicus
7	G207	286661	1.20E-36	Oryza sativa
9	G241	6552360	2.60E-54	Nicotiana tabacum
9	G241	6782745	2.20E-53	Oryza sativa
9	G241	8097368	5.70E-53	Hordeum vulgare
9	G241	20560	1.80E-52	Petunia x hybrida
9	G241	7217727	2.70E-52	Sorghum bicolor
9	G241	5891408	4.60E-52	Lycopersicon esculentum
9	G241	5139803	7.40E-52	Glycine max
9	G241	7560175	4.10E-50	Medicago truncatula
9	G241	8381332	1.40E-44	Gossypium arboreum
9	G241	4886263	1.20E-42	Antirrhinum majus
11	G254	5847380	2.00E-41	Zea mays
11	G254	7614730	2.90E-41	Lotus japonicus

Figure 3B

SEQ ID No.	GID	Genbank NID	P-value	Species
11	G254	9204594	4.80E-41	Glycine max
11	G254	9193761	6.70E-37	Medicago truncatula
11	G254	6907081	1.40E-35	Oryza sativa
11	G254	6976741	4.30E-33	Lycopersicon esculentum
11	G254	8903196	4.20E-31	Hordeum vulgare
11	G254	9424828	3.50E-25	Triticum aestivum
11	G254	6858452	3.40E-23	Sorghum bicolor
11	G254	3003284	0.00068	Mesembryanthemum crystallinum
13	G263	5821135	1.70E-73	Nicotiana tabacum
13	G263	19487	7.90E-69	Lycopersicon peruvianum
13	G263	662929	5.30E-65	Glycine max
13	G263	7766273	9.20E-49	Medicago truncatula
13	G263	7720908	3.60E-42	Lotus japonicus
13	G263	9303509	2.40E-37	Sorghum bicolor
13	G263	3326480	2.20E-36	Gossypium hirsutum
13	G263	8107182	5.10E-35	Lycopersicon esculentum
13	G263	8381330	7.00E-34	Gossypium arboreum
13	G263	4528238	6.60E-29	Citrus unshiu
15	G308	5640156	3.50E-162	Triticum aestivum
15	G308	5640154	2.30E-134	Zea mays
15	G308	6970471	4.20E-120	Oryza sativa
15	G308	7718432	8.70E-80	Medicago truncatula
15	G308	8330344	3.90E-76	Mesembryanthemum crystallinum
15	G308	5047560	1.50E-71	Gossypium hirsutum
15	G308	7588689	1.90E-68	Glycine max
15	G308	7623983	2.90E-62	Gossypium arboreum
15	G308	7780253	1.10E-57	Lotus japonicus
15	G308	6733213	3.70E-48	Lycopersicon esculentum
17	G536	2689478	9.50E-69	Nicotiana tabacum
17	G536	1773327	4.60E-68	Mesembryanthemum crystallinum
17	G536	2921511	5.30E-68	Fritillaria agrestis
17	G536	1575724	7.30E-68	Glycine max
17	G536	8515887	9.20E-68	Populus alba x Populus tremula
17	G536	6179980	1.70E-67	Lilium longiflorum
17	G536	1519250	8.60E-67	Oryza sativa
17	G536	1321992	4.30E-66	Solanum tuberosum
17	G536	7535681	9.50E-66	Sorghum bicolor
17	G536	555973	1.30E-65	Pisum sativum
19	G680	9258166	5.70E-36	Glycine max
19	G680	9255178	3.00E-29	Zea mays
19	G680	5274804	1.20E-27	Lycopersicon esculentum
19	G680	4974199	3.00E-22	Oryza sativa
19	G680	3325786	2.10E-21	Gossypium hirsutum
19	G680	9119112	1.30E-18	Medicago truncatula
19	G680	7660673	3.20E-17	Sorghum bicolor
19	G680	7243970	6.10E-16	Mentha x piperita
19	G680	3858093	2.10E-10	Populus balsamifera subsp. trichocarpa
19	G680	8845091	3.70E-10	Triticum aestivum
21	G867	7643366	1.80E-57	Medicago truncatula
21	G867	8329389	9.00E-51	Mesembryanthemum crystallinum
21	G867	8669779	2.20E-46	Glycine max
21	G867	9430646	5.70E-40	Lycopersicon esculentum
21	G867	8902194	1.20E-34	Hordeum vulgare

Figure 3C

SEQ ID No.	GID	Genbank NID	P-value	Species
21	G867	7722547	1.00E-33	Lotus japonicus
21	G867	7324245	3.10E-32	Lycopersicon pennellii
21	G867	8749037	1.10E-31	Citrus x paradisi
21	G867	6069643	2.50E-29	Oryza sativa
21	G867	9302986	1.40E-28	Sorghum bicolor
23	G912	5616085	8.60E-71	Brassica napus
23	G912	7410271	5.70E-46	Lycopersicon esculentum
23	G912	7719106	5.20E-43	Medicago truncatula
23	G912	6667103	2.30E-38	Glycine max
23	G912	6983854	1.30E-34	Oryza sativa
23	G912	7324530	1.00E-32	Lycopersicon pennellii
23	G912	8904571	9.20E-29	Triticum aestivum
23	G912	7740143	1.90E-28	Lotus japonicus
23	G912	7644788	2.10E-19	Pinus taeda
23	G912	5050536	9.20E-18	Gossypium hirsutum
25	G996	7566043	2.30E-65	Medicago truncatula
25	G996	7535969	1.00E-61	Sorghum bicolor
25	G996	7339127	1.80E-59	Lycopersicon esculentum
25	G996	6341619	5.60E-59	Glycine max
25	G996	8381332	7.20E-43	Gossypium arboreum
25	G996	5049507	5.00E-41	Gossypium hirsutum
25	G996	6850206	2.10E-40	Oryza sativa
25	G996	7776223	2.20E-40	Lotus japonicus
25	G996	19058	5.30E-39	Hordeum vulgare
25	G996	4680189	6.00E-39	Oryza sativa subsp. indica
27	G1068	7333976	1.70E-27	Lycopersicon esculentum
27	G1068	4405544	3.20E-27	Glycine max
27	G1068	7009437	5.50E-23	Zea mays
27	G1068	7536402	5.80E-23	Sorghum bicolor
27	G1068	3107210	7.20E-21	Oryza sativa
27	G1068	3819186	5.80E-18	Hordeum vulgare
27	G1068	7624850	8.40E-18	Gossypium arboreum
27	G1068	9411568	1.90E-13	Triticum aestivum
27	G1068	5419913	3.50E-13	Lactuca sativa
27	G1068	7721066	8.90E-13	Lotus japonicus
29	G1337	7410432	2.60E-41	Lycopersicon esculentum
29	G1337	3618319	1.10E-32	Oryza sativa
29	G1337	7571599	1.00E-28	Medicago truncatula
29	G1337	7685955	5.10E-27	Glycine max
29	G1337	7323708	2.60E-25	Lycopersicon hirsutum
29	G1337	4091805	1.00E-18	Malus domestica
29	G1337	6917805	4.80E-18	Lycopersicon pennellii
29	G1337	3341722	1.60E-17	Raphanus sativus
29	G1337	2303680	4.50E-17	Brassica napus
29	G1337	4557092	9.10E-17	Pinus radiata
31	G1141	8346772	9.90E-46	Catharanthus roseus
31	G1141	7205636	3.60E-40	Medicago truncatula
31	G1141	7590901	5.40E-40	Glycine max
31	G1141	7777379	6.10E-38	Lotus japonicus
31	G1141	9363798	8.00E-36	Triticum aestivum
31	G1141	8903111	6.10E-31	Hordeum vulgare
31	G1141	568076	1.00E-23	Oryza sativa
31	G1141	6527472	1.10E-17	Lycopersicon esculentum

Figure 3D

SEQ ID No.	GID	Genbank NID	P-value	Species
31	G1141	7324705	1.70E-16	Lycopersicon pennellii
31	G1141	7624302	1.80E-16	Gossypium arboreum
33	G46	5760554	4.00E-29	Glycine max
33	G46	7778996	4.20E-28	Lotus japonicus
33	G46	5050094	1.70E-26	Gossypium hirsutum
33	G46	790361	3.60E-26	Nicotiana tabacum
33	G46	5603736	7.30E-24	Lycopersicon esculentum
33	G46	7238955	1.20E-23	Medicago truncatula
33	G46	8809574	4.10E-23	Nicotiana sylvestris
33	G46	7528275	1.40E-22	Mesembryanthemum crystallinum
33	G46	8980312	1.60E-22	Catharanthus roseus
33	G46	6478844	2.40E-22	Matricaria chamomilla
35	G242	6529807	1.90E-70	Lycopersicon esculentum
35	G242	7624453	3.00E-63	Gossypium arboreum
35	G242	7564212	2.30E-62	Medicago truncatula
35	G242	5820271	3.70E-60	Glycine max
35	G242	7322467	1.10E-55	Lycopersicon hirsutum
35	G242	5045349	1.80E-51	Gossypium hirsutum
35	G242	8071527	6.80E-46	Solanum tuberosum
35	G242	7790004	7.40E-45	Beta vulgaris
35	G242	7746594	4.70E-41	Lotus japonicus
35	G242	286661	3.40E-39	Oryza sativa
37	G227	6529807	4.80E-67	Lycopersicon esculentum
37	G227	7624453	2.50E-66	Gossypium arboreum
37	G227	5045349	7.90E-65	Gossypium hirsutum
37	G227	7322467	8.00E-60	Lycopersicon hirsutum
37	G227	5820271	2.60E-59	Glycine max
37	G227	9199531	4.70E-57	Medicago truncatula
37	G227	8071527	9.30E-49	Solanum tuberosum
37	G227	7790004	8.30E-46	Beta vulgaris
37	G227	7746594	1.70E-45	Lotus japonicus
37	G227	286661	9.20E-37	Oryza sativa
39	G1307	8172759	2.90E-56	Medicago truncatula
39	G1307	5139807	1.60E-54	Glycine max
39	G1307	1370139	2.20E-47	Lycopersicon esculentum
39	G1307	1946264	6.20E-45	Oryza sativa
39	G1307	6552360	6.50E-45	Nicotiana tabacum
39	G1307	7500978	4.60E-39	Gossypium arboreum
39	G1307	7217727	8.70E-36	Sorghum bicolor
39	G1307	7746498	9.40E-34	Lotus japonicus
39	G1307	517491	9.90E-34	Zea mays
39	G1307	8097368	1.70E-33	Hordeum vulgare
41	G1327	5139803	1.10E-44	Glycine max
41	G1327	7560175	1.20E-44	Medicago truncatula
41	G1327	6782745	6.60E-44	Oryza sativa
41	G1327	5891408	2.30E-43	Lycopersicon esculentum
41	G1327	7217727	3.10E-43	Sorghum bicolor
41	G1327	20560	2.40E-41	Petunia x hybrida
41	G1327	6552360	5.40E-40	Nicotiana tabacum
41	G1327	8097368	9.80E-40	Hordeum vulgare
41	G1327	8381332	5.20E-39	Gossypium arboreum
41	G1327	9252441	1.60E-38	Solanum tuberosum
43	G673	6062169	4.90E-36	Lycopersicon esculentum

Figure 3E

SEQ ID No.	GID	Genbank NID	P-value	Species
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43	G673	9205170	2.60E-28	Glycine max
43	G673	5847380	7.10E-26	Zea mays
43	G673	7614730	7.70E-26	Lotus japonicus
43	G673	9193761	3.40E-25	Medicago truncatula
43	G673	9424828	1.70E-24	Triticum aestivum
43	G673	8903196	3.40E-23	Hordeum vulgare
43	G673	6858452	2.80E-14	Sorghum bicolor
43	G673	3003284	1.50E-09	Mesembryanthemum crystallinum
45	G307	5640156	3.80E-151	Triticum aestivum
45	G307	5640154	1.00E-101	Zea mays
45	G307	6970471	1.70E-97	Oryza sativa
45	G307	7718432	4.00E-82	Medicago truncatula
45	G307	8330344	7.90E-78	Mesembryanthemum crystallinum
45	G307	5047560	1.00E-72	Gossypium hirsutum
45	G307	7588689	2.70E-69	Glycine max
45	G307	7623983	2.20E-64	Gossypium arboreum
45	G307	7780253	9.30E-59	Lotus japonicus
45	G307	6733213	1.90E-51	Lycopersicon esculentum
47	G529	1773327	8.80E-117	Mesembryanthemum crystallinum
47	G529	8515887	1.20E-115	Populus alba x Populus tremula
47	G529	6179980	2.30E-115	Lilium longiflorum
47	G529	2921511	2.40E-115	Fritillaria agrestis
47	G529	1575724	3.30E-115	Glycine max
47	G529	466335	2.80E-112	Lycopersicon esculentum
47	G529	1519250	4.10E-112	Oryza sativa
47	G529	2689478	2.20E-110	Nicotiana tabacum
47	G529	2266661	5.10E-109	Hordeum vulgare
47	G529	1321992	1.20E-108	Solanum tuberosum
49	G531	2921511	7.40E-109	Fritillaria agrestis
49	G531	6179980	2.30E-108	Lilium longiflorum
49	G531	1773327	4.50E-108	Mesembryanthemum crystallinum
49	G531	8515887	8.90E-108	Populus alba x Populus tremula
49	G531	2689478	2.10E-107	Nicotiana tabacum
49	G531	1575724	4.90E-107	Glycine max
49	G531	1519250	1.70E-106	Oryza sativa
49	G531	466335	1.20E-104	Lycopersicon esculentum
49	G531	1321992	1.10E-103	Solanum tuberosum
49	G531	2266661	1.60E-103	Hordeum vulgare
51	G214	8170933	8.80E-35	Lycopersicon esculentum
51	G214	9205339	1.20E-27	Glycine max
51	G214	8577344	1.80E-23	Zea mays
51	G214	9119112	2.40E-18	Medicago truncatula
51	G214	7660673	4.80E-15	Sorghum bicolor
51	G214	8213273	4.40E-14	Oryza sativa
51	G214	3325786	4.70E-10	Gossypium hirsutum
51	G214	9435251	1.50E-09	Hordeum vulgare
51	G214	9411569	6.80E-09	Triticum aestivum
51	G214	7614730	3.00E-07	Lotus japonicus
53	G1930	7643366	7.70E-57	Medicago truncatula
53	G1930	8329389	3.60E-47	Mesembryanthemum crystallinum
53	G1930	6069592	8.60E-47	Glycine max
53	G1930	9430646	6.60E-39	Lycopersicon esculentum

Figure 3F

SEQ ID No.	GID	Genbank NID	P-value	Species
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53	G1930	7324245	9.20E-33	Lycopersicon pennellii
53	G1930	8902194	2.40E-31	Hordeum vulgare
53	G1930	9247126	5.90E-28	Oryza sativa
53	G1930	8749037	2.40E-27	Citrus x paradisi
53	G1930	9302986	6.40E-26	Sorghum bicolor
55	G9	7643366	5.40E-56	Medicago truncatula
55	G9	8669779	3.30E-50	Glycine max
55	G9	8329389	1.20E-48	Mesembryanthemum crystallinum
55	G9	7412012	1.20E-41	Lycopersicon esculentum
55	G9	8902194	6.60E-36	Hordeum vulgare
55	G9	7722547	2.10E-33	Lotus japonicus
55	G9	7324245	1.90E-32	Lycopersicon pennellii
55	G9	8749037	1.10E-31	Citrus x paradisi
55	G9	9247126	1.20E-29	Oryza sativa
55	G9	9302986	7.00E-29	Sorghum bicolor
57	G993	7643366	9.50E-59	Medicago truncatula
57	G993	8329389	8.10E-50	Mesembryanthemum crystallinum
57	G993	8669779	4.80E-49	Glycine max
57	G993	4384549	4.20E-40	Lycopersicon esculentum
57	G993	8902194	2.00E-34	Hordeum vulgare
57	G993	7719409	1.00E-32	Lotus japonicus
57	G993	8749037	4.10E-32	Citrus x paradisi
57	G993	9247126	1.00E-30	Oryza sativa
57	G993	7324245	1.20E-30	Lycopersicon pennellii
57	G993	9302986	9.10E-27	Sorghum bicolor
59	G41	5616085	6.30E-84	Brassica napus
59	G41	5603726	2.60E-50	Lycopersicon esculentum
59	G41	7719106	2.00E-43	Medicago truncatula
59	G41	6667103	1.60E-37	Glycine max
59	G41	6983854	1.80E-33	Oryza sativa
59	G41	7324530	9.50E-30	Lycopersicon pennellii
59	G41	8904571	2.70E-29	Triticum aestivum
59	G41	7740143	2.50E-26	Lotus japonicus
59	G41	7644788	3.40E-19	Pinus taeda
59	G41	7625186	6.50E-19	Gossypium arboreum
61	G40	5616085	7.70E-86	Brassica napus
61	G40	5603726	1.60E-50	Lycopersicon esculentum
61	G40	7719106	4.70E-42	Medicago truncatula
61	G40	6667103	1.10E-36	Glycine max
61	G40	6983854	4.70E-35	Oryza sativa
61	G40	8904571	3.50E-29	Triticum aestivum
61	G40	7324530	5.20E-29	Lycopersicon pennellii
61	G40	7740143	1.40E-25	Lotus japonicus
61	G40	7644788	1.80E-20	Pinus taeda
61	G40	7625186	5.70E-20	Gossypium arboreum
63	G42	5616085	8.60E-87	Brassica napus
63	G42	5603726	2.20E-53	Lycopersicon esculentum
63	G42	7719106	5.20E-43	Medicago truncatula
63	G42	6667103	6.00E-38	Glycine max
63	G42	6983854	1.10E-35	Oryza sativa
63	G42	8904571	5.50E-31	Triticum aestivum

Figure 3G

SEQ ID No.	GID	Genbank NID	P-value	Species
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63	G42	7644788	2.40E-20	Pinus taeda
63	G42	7625186	1.50E-19	Gossypium arboreum
65	G1127	6913305	2.60E-29	Glycine max
65	G1127	9280727	5.40E-27	Oryza sativa
65	G1127	2213533	7.00E-24	Pisum sativum
65	G1127	7009437	4.70E-23	Zea mays
65	G1127	7536402	5.00E-23	Sorghum bicolor
65	G1127	7333976	1.20E-20	Lycopersicon esculentum
65	G1127	3819186	6.20E-16	Hordeum vulgare
65	G1127	7624850	1.60E-15	Gossypium arboreum
65	G1127	4165182	2.80E-12	Antirrhinum majus
65	G1127	7765939	5.10E-09	Medicago truncatula
67	G2657	7238733	2.70E-66	Medicago truncatula
67	G2657	6846994	7.60E-55	Glycine max
67	G2657	7615218	1.10E-43	Lotus japonicus
67	G2657	9445090	4.00E-41	Triticum aestivum
67	G2657	7333102	3.20E-38	Lycopersicon esculentum
67	G2657	9252370	1.90E-27	Solanum tuberosum
67	G2657	5042437	5.90E-21	Oryza sativa
67	G2657	7536402	8.60E-20	Sorghum bicolor
67	G2657	7624850	2.20E-18	Gossypium arboreum
67	G2657	7009437	1.80E-16	Zea mays
69	G326	7410432	1.10E-37	Lycopersicon esculentum
69	G326	3618319	2.90E-32	Oryza sativa
69	G326	7571599	4.90E-30	Medicago truncatula
69	G326	7232283	6.30E-28	Glycine max
69	G326	7323708	6.00E-27	Lycopersicon hirsutum
69	G326	4091805	2.30E-19	Malus domestica
69	G326	6917805	6.50E-19	Lycopersicon pennellii
69	G326	3341722	2.50E-18	Raphanus sativus
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mbi19 Sequence Listing.ST25
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mbi19 Sequence Listing.ST25

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mbil9 Sequence Listing.ST25

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Ala Glu Arg Leu Lys Arg Trp Lys Glu Tyr Asn Glu Thr Val Glu Glu
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Val Ser Thr Lys Lys Arg Lys Val Pro Ala Lys Gly Ser Lys Lys Gly
      50                      55                      60

Cys Met Lys Gly Lys Gly Gly Pro Glu Asn Ser Arg Cys Ser Phe Arg
      65                      70                      75                      80

Gly Val Arg Gln Arg Ile Trp Gly Lys Trp Val Ala Glu Ile Arg Glu
      85                      90                      95

Pro Asn Arg Gly Ser Arg Leu Trp Leu Gly Thr Phe Pro Thr Ala Gln
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Glu Ala Ala Ser Ala Tyr Asp Glu Ala Ala Lys Ala Met Tyr Gly Pro
      115                     120                     125

Leu Ala Arg Leu Asn Phe Pro Arg Ser Asp Ala Ser Glu Val Thr Ser
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Thr Ser Ser Gln Ser Glu Val Cys Thr Val Glu Thr Pro Gly Cys Val
      145                     150                     155                     160

His Val Lys Thr Glu Asp Pro Asp Cys Glu Ser Lys Pro Phe Ser Gly
      165                     170                     175

Gly Val Glu Pro Met Tyr Cys Leu Glu Asn Gly Ala Glu Glu Met Lys
      180                     185                     190

Arg Gly Val Lys Ala Asp Lys His Trp Leu Ser Glu Phe Glu His Asn
      195                     200                     205

Tyr Trp Ser Asp Ile Leu Lys Glu Lys Glu Lys Gln Lys Glu Gln Gly
      210                     215                     220

Ile Val Glu Thr Cys Gln Gln Gln Gln Gln Asp Ser Leu Ser Val Ala
      225                     230                     235                     240

Asp Tyr Gly Trp Pro Asn Asp Val Asp Gln Ser His Leu Asp Ser Ser
      245                     250                     255

Asp Met Phe Asp Val Asp Glu Leu Leu Arg Asp Leu Asn Gly Asp Asp
      260                     265                     270

Val Phe Ala Gly Leu Asn Gln Asp Arg Tyr Pro Gly Asn Ser Val Ala
      275                     280                     285

Asn Gly Ser Tyr Arg Pro Glu Ser Gln Gln Ser Gly Phe Asp Pro Leu
      290                     295                     300

Gln Ser Leu Asn Tyr Gly Ile Pro Pro Phe Gln Leu Glu Gly Lys Asp
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aaaaaaaaa aaaaaaaaaa aaaaaaaaaa aaaaaaaaaa aaaaaa 909

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35 40 45

Asn Pro Val Pro Lys Leu Glu Pro Ser Ser Pro Val Leu Asp Pro Asp
50 55 60

Ser Tyr Val Gln Glu Ile Leu Gln Met Glu Ala Glu Ser Ser Ser Ser
65 70 75 80

Ser Ser Thr Thr Thr Ser Pro Glu Val Glu Thr Val Ser Asn Arg Lys
85 90 95

Lys Thr Lys Arg Phe Glu Glu Thr Arg His Tyr Arg Gly Val Arg Arg
100 105 110

Arg Pro Trp Gly Lys Phe Ala Ala Glu Ile Arg Asp Pro Ala Lys Lys
115 120 125

Gly Ser Arg Ile Trp Leu Gly Thr Phe Glu Ser Asp Ile Asp Ala Ala
130 135 140

Arg Ala Tyr Asp Tyr Ala Ala Phe Lys Leu Arg Gly Arg Lys Ala Val
145 150 155 160

Leu Asn Phe Pro Leu Asp Ala Gly Lys Tyr Asp Ala Pro Val Asn Ser
165 170 175

Cys Arg Lys Arg Arg Arg Thr Asp Val Pro Gln Pro Gln Gly Thr Thr
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Thr Ser Thr Ser Ser Ser Ser Ser Asn
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mbi19 Sequence Listing.ST25

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Glu Asp Glu Gln Leu Arg Arg Met Val Glu Lys Tyr Gly Pro Arg Asn	
15 20 25	
 tgg tct gcg att agc aaa tcg att cca ggt cga tct ggt aaa tcg tgt	147
Trp Ser Ala Ile Ser Lys Ser Ile Pro Gly Arg Ser Gly Lys Ser Cys	
30 35 40	
 aga tta cgt tgg tgt aat cag tta tct ccg gag gtt gag cat cgt cct	195
Arg Leu Arg Trp Cys Asn Gln Leu Ser Pro Glu Val Glu His Arg Pro	
45 50 55 60	
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Phe Ser Pro Glu Glu Asp Glu Thr Ile Val Thr Ala Arg Ala Gln Phe	
65 70 75	
 ggg aac aag tgg gcg acg att gct cgt ctt ctt aac ggt cgt acg gat	291
Gly Asn Lys Trp Ala Thr Ile Ala Arg Leu Leu Asn Gly Arg Thr Asp	
80 85 90	
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Asn Ala Val Lys Asn His Trp Asn Ser Thr Leu Lys Arg Lys Cys Ser	
95 100 105	
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Gly Gly Val Ala Val Thr Thr Val Thr Glu Thr Glu Glu Asp Gln Asp	
110 115 120	
 cgg ccg aag aag agg aga tct gtt agc ttt gat cct gct ttt gct ccg	435
Arg Pro Lys Lys Arg Arg Ser Val Ser Phe Asp Pro Ala Phe Ala Pro	
125 130 135 140	
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Val Asp Thr Gly Leu Tyr Met Ser Pro Glu Ser Pro Asn Gly Ile Asp	
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Val Ser Asp Ser Ser Thr Ile Pro Ser Pro Ser Ser Pro Val Ala Gln	
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Leu Phe Lys Pro Met Pro Ile Ser Gly Gly Phe Thr Val Val Pro Gln	
175 180 185	
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Pro Leu Pro Val Glu Met Ser Ser Ser Ser Glu Asp Pro Pro Thr Ser	
190 195 200	
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Leu Ser Leu Ser Leu Pro Gly Ala Glu Asn Thr Ser Ser Ser His Asn	
205 210 215 220	
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Asn Asn Asn Asn Ala Leu Met Phe Pro Arg Phe Glu Ser Gln Met Lys	
225 230 235	
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Ile Asn Val Glu Glu Arg Gly Gly Gly Gly Glu Gly Arg Arg Gly Glu	
240 245 250	
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Phe Met Thr Val Val Gln Glu Met Ile Lys Ala Glu Val Arg Ser Tyr	
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Met Ala Glu Met Gln Lys Thr Ser Gly Gly Phe Val Val Gly Gly Leu	
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mbi19 Sequence Listing.ST25

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 Tyr Glu Ser Gly Gly Asn Gly Gly Phe Arg Asp Cys Gly Val Ile Thr
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cct aag gtt gag tag ttttggttta gggttaaaac ttgaatcgat tggggatttt 970
 Pro Lys Val Glu

caagagcatt cttttttggg gtttatggta aaattaaaaa caaaaacaaa atgtacagag 1030

gaattaaaaat ttctatggaa taatcttaaa tctcaaatat ttgttacttg ttttggtgat 1090

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 35 40 45

Cys Asn Gln Leu Ser Pro Glu Val Glu His Arg Pro Phe Ser Pro Glu
 50 55 60

Glu Asp Glu Thr Ile Val Thr Ala Arg Ala Gln Phe Gly Asn Lys Trp
 65 70 75 80

Ala Thr Ile Ala Arg Leu Leu Asn Gly Arg Thr Asp Asn Ala Val Lys
 85 90 95

Asn His Trp Asn Ser Thr Leu Lys Arg Lys Cys Ser Gly Gly Val Ala
 100 105 110

Val Thr Thr Val Thr Glu Thr Glu Glu Asp Gln Asp Arg Pro Lys Lys
 115 120 125

Arg Arg Ser Val Ser Phe Asp Pro Ala Phe Ala Pro Val Asp Thr Gly
 130 135 140

Leu Tyr Met Ser Pro Glu Ser Pro Asn Gly Ile Asp Val Ser Asp Ser
 145 150 155 160

Ser Thr Ile Pro Ser Pro Ser Ser Pro Val Ala Gln Leu Phe Lys Pro
 165 170 175

Met Pro Ile Ser Gly Gly Phe Thr Val Val Pro Gln Pro Leu Pro Val
 180 185 190

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210 215 220

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Pro Cys Cys Glu Lys Met Gly Leu Lys Arg Gly Pro Trp Thr Pro Glu																
5 10 15 20																
gaa gat caa atc ttg gtc tct ttt atc ctc aac cat gga cat agt aac															153	
Glu Asp Gln Ile Leu Val Ser Phe Ile Leu Asn His Gly His Ser Asn																
25 30 35																
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Trp Arg Ala Leu Pro Lys Gln Ala Gly Leu Leu Arg Cys Gly Lys Ser																
40 45 50																
tgt aga ctt agg tgg atg aac tat tta aag cct gat att aaa cgt ggc															249	
Cys Arg Leu Arg Trp Met Asn Tyr Leu Lys Pro Asp Ile Lys Arg Gly																
55 60 65																
aat ttc acc aaa gaa gag gaa gat gct atc atc agc tta cac caa ata															297	
Asn Phe Thr Lys Glu Glu Glu Asp Ala Ile Ile Ser Leu His Gln Ile																
70 75 80																
ctt ggc aat aga tgg tca gcg att gca gca aaa ctg cct gga aga acc															345	
Leu Gly Asn Arg Trp Ser Ala Ile Ala Ala Lys Leu Pro Gly Arg Thr																
85 90 95 100																
gat aac gag atc aag aac gta tgg cac act cac ttg aag aag aga ctc															393	
Asp Asn Glu Ile Lys Asn Val Trp His Thr His Leu Lys Lys Arg Leu																
105 110 115																
gaa gat tat caa cca gct aaa cct aag acc agc aac aaa aag aag ggt															441	
Glu Asp Tyr Gln Pro Ala Lys Pro Lys Thr Ser Asn Lys Lys Lys Gly																
120 125 130																
act aaa cca aaa tct gaa tcc gta ata acg agc tcg aac agt act aga															489	
Thr Lys Pro Lys Ser Glu Ser Val Ile Thr Ser Ser Asn Ser Thr Arg																
135 140 145																

mbi19 Sequence Listing.ST25

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Phe Ser Thr Ser Pro Ser Thr Ser Glu Val Ser Ser Met Thr Leu Ile
    165                      170                      175                      180

agc cac gac ggc tat agc aac gag att aat atg gat aac aaa ccg gga      633
Ser His Asp Gly Tyr Ser Asn Glu Ile Asn Met Asp Asn Lys Pro Gly
    185                      190                      195

gat atc agt act atc gat caa gaa tgt gtt tct ttc gaa act ttt ggt      681
Asp Ile Ser Thr Ile Asp Gln Glu Cys Val Ser Phe Glu Thr Phe Gly
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gcg gat atc gat gaa agc ttc tgg aaa gag aca ctg tat agc caa gat      729
Ala Asp Ile Asp Glu Ser Phe Trp Lys Glu Thr Leu Tyr Ser Gln Asp
    215                      220                      225

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Glu His Asn Tyr Val Ser Asn Asp Leu Glu Val Ala Gly Leu Val Glu
    230                      235                      240

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    245                      250                      255                      260

ttt gac agt gag atg gaa ctt ctg gtt cga tgt att ggc tag              867
Phe Asp Ser Glu Met Glu Leu Leu Val Arg Cys Ile Gly
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gaaatggtgc aaattagtta aggctaagaa attcaaaagc ttttgtttac cgagaaaaaa      987

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Gly His Ser Asn Trp Arg Ala Leu Pro Lys Gln Ala Gly Leu Leu Arg
35      40      45

Cys Gly Lys Ser Cys Arg Leu Arg Trp Met Asn Tyr Leu Lys Pro Asp
50      55      60

Ile Lys Arg Gly Asn Phe Thr Lys Glu Glu Glu Asp Ala Ile Ile Ser
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Leu His Gln Ile Leu Gly Asn Arg Trp Ser Ala Ile Ala Ala Lys Leu
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Pro Gly Arg Thr Asp Asn Glu Ile Lys Asn Val Trp His Thr His Leu
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Lys Lys Arg Leu Glu Asp Tyr Gln Pro Ala Lys Pro Lys Thr Ser Asn

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mbi19 Sequence Listing.ST25

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145	150	155	160
Gly Glu Ser Leu Phe Ser Thr Ser Pro Ser Thr Ser Glu Val Ser Ser			
165	170	175	
Met Thr Leu Ile Ser His Asp Gly Tyr Ser Asn Glu Ile Asn Met Asp			
180	185	190	
Asn Lys Pro Gly Asp Ile Ser Thr Ile Asp Gln Glu Cys Val Ser Phe			
195	200	205	
Glu Thr Phe Gly Ala Asp Ile Asp Glu Ser Phe Trp Lys Glu Thr Leu			
210	215	220	
Tyr Ser Gln Asp Glu His Asn Tyr Val Ser Asn Asp Leu Glu Val Ala			
225	230	235	240
Gly Leu Val Glu Ile Gln Gln Glu Phe Gln Asn Leu Gly Ser Ala Asn			
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Ser Ile Pro Ala Thr Gly Arg Thr Ser Thr Val Ser Phe Ser Glu Asp		
30 35 40		
ccg acg acg aag att cgg aag ccg tac aca atc aag aag tcg aga gag		194
Pro Thr Thr Lys Ile Arg Lys Pro Tyr Thr Ile Lys Lys Ser Arg Glu		
45 50 55 60		
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Asn Trp Thr Asp Gln Glu His Asp Lys Phe Leu Glu Ala Leu His Leu		
65 70 75		

mbi19 Sequence Listing.ST25

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95	100 105	
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175	180 185	
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190	195 200	
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205	210 215 220	
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225	230 235	
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240	245 250	
tca ggc cac ctc cag aga tta aag cag atg gat cca ata aat atg gaa Ser Gly His Leu Gln Arg Leu Lys Gln Met Asp Pro Ile Asn Met Glu	818	
255	260 265	
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270	275 280	
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285	290 295 300	
ttg aaa tag agatagaata aaacaataat gtaccttatg tgagatcaag Leu Lys	963	
agacaatcat ccaaggtctg tatgcattgc ttggatttag gcctcgtgtt ctactacag	1023	
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mbi19 Sequence Listing.ST25

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 35 40 45

Ile Arg Lys Pro Tyr Thr Ile Lys Lys Ser Arg Glu Asn Trp Thr Asp
 50 55 60

Gln Glu His Asp Lys Phe Leu Glu Ala Leu His Leu Phe Asp Arg Asp
 65 70 75 80

Trp Lys Lys Ile Glu Ala Phe Val Gly Ser Lys Thr Val Val Gln Ile
 85 90 95

Arg Ser His Ala Gln Lys Tyr Phe Leu Lys Val Gln Lys Ser Gly Ala
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Asn Glu His Leu Pro Leu Pro Arg Pro Lys Arg Lys Ala Ser His Pro
 115 120 125

Tyr Pro Ile Lys Ala Pro Lys Asn Val Ala Tyr Thr Ser Leu Pro Ser
 130 135 140

Ser Ser Thr Leu Pro Leu Leu Glu Pro Gly Tyr Leu Tyr Ser Ser Asp
 145 150 155 160

Ser Lys Ser Leu Met Gly Asn Gln Ala Val Cys Ala Ser Thr Ser Ser
 165 170 175

Ser Trp Asn His Glu Ser Thr Asn Leu Pro Lys Pro Val Ile Glu Glu
 180 185 190

Glu Pro Gly Val Ser Ala Thr Ala Pro Leu Pro Asn Asn Arg Cys Arg
 195 200 205

Gln Glu Asp Thr Glu Arg Val Arg Ala Val Thr Lys Pro Asn Asn Glu
 210 215 220

Glu Ser Cys Glu Lys Pro His Arg Val Met Pro Asn Phe Ala Glu Val
 225 230 235 240

Tyr Ser Phe Ile Gly Ser Val Phe Asp Pro Asn Thr Ser Gly His Leu
 245 250 255

Gln Arg Leu Lys Gln Met Asp Pro Ile Asn Met Glu Thr Val Leu Leu
 260 265 270

mbil9 Sequence Listing.ST25

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 <223> G263

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 Val Thr Ala Ala Gln Arg Ser Val Pro Ala Pro Phe Leu Ser Lys Thr
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 Tyr Gln Leu Val Asp Asp His Ser Thr Asp Asp Val Val Ser Trp Asn
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 Glu Glu Gly Thr Ala Phe Val Val Trp Lys Thr Ala Glu Phe Ala Lys
 40 45 50

gat ctt ctt cct caa tac ttc aag cat aat aat ttc tca agc ttc att 248
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 55 60 65

cgt cag ctc aac act tac gga ttt cgt aaa act gta ccg gat aaa tgg 296
 Arg Gln Leu Asn Thr Tyr Gly Phe Arg Lys Thr Val Pro Asp Lys Trp
 70 75 80

gaa ttt gca aac gat tat ttc cgg aga ggc ggg gag gat ctg ttg acg 344
 Glu Phe Ala Asn Asp Tyr Phe Arg Arg Gly Gly Glu Asp Leu Leu Thr
 85 90 95

gac ata cga cgg cgt aaa tcg gtg att gct tca acg gcg ggg aaa tgt 392
 Asp Ile Arg Arg Arg Lys Ser Val Ile Ala Ser Thr Ala Gly Lys Cys
 100 105 110 115

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 Val Val Val Gly Ser Pro Ser Glu Ser Asn Ser Gly Gly Gly Asp Asp
 120 125 130

cac ggt tca agc tcc acg tca tca ccc ggt tcg tcg aag aat cct ggt 488
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 135 140 145

tcg gtg gag aac atg gtt gct gat tta tca gga gag aac gag aag ctt 536
 Ser Val Glu Asn Met Val Ala Asp Leu Ser Gly Glu Asn Glu Lys Leu
 150 155 160

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 165 170 175

cag cgc gat gag cta gtg acg ttc ttg acg ggt cat ctg aaa gta aga 632
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mbil9 Sequence Listing.ST25

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 215 220 225

gca gag gag ggg gta ggt gaa gga ttg aaa ttg ttt ggg gtg tgg ttg 776
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 230 235 240

aaa gga gag aga aaa aag agg gac cgg gat gaa aag aat tat gtg gtg 824
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 245 250 255

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 Ser Gly Ser Arg Met Thr Glu Ile Lys Asn Val Asp Phe His Ala Pro
 260 265 270 275

ttg tgg aaa agc agc aaa gtc tgc aac taa aaaaagagta gaagactggt 922
 Leu Trp Lys Ser Ser Lys Val Cys Asn
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caaaccagcg tgtgacacgt catcgacgac gacgaaaaaa atgatttaaa aaactatttt 982

tttccgtaag gaagaaaagt tatttttatg ttttaaaaag gtgaagaagg tccagaagga 1042

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 35 40 45

Phe Ala Lys Asp Leu Leu Pro Gln Tyr Phe Lys His Asn Asn Phe Ser
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Ser Phe Ile Arg Gln Leu Asn Thr Tyr Gly Phe Arg Lys Thr Val Pro
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Asp Lys Trp Glu Phe Ala Asn Asp Tyr Phe Arg Arg Gly Gly Glu Asp
 85 90 95

Leu Leu Thr Asp Ile Arg Arg Arg Lys Ser Val Ile Ala Ser Thr Ala
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Gly Lys Cys Val Val Val Gly Ser Pro Ser Glu Ser Asn Ser Gly Gly
 115 120 125

Gly Asp Asp His Gly Ser Ser Ser Thr Ser Ser Pro Gly Ser Ser Lys
 130 135 140

mbi19 Sequence Listing.ST25

Asn Pro Gly Ser Val Glu Asn Met Val Ala Asp Leu Ser Gly Glu Asn
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Glu Lys Leu Lys Arg Glu Asn Asn Asn Leu Ser Ser Glu Leu Ala Ala
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Ala Lys Lys Gln Arg Asp Glu Leu Val Thr Phe Leu Thr Gly His Leu
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Lys Val Arg Pro Glu Gln Ile Asp Lys Met Ile Lys Gly Gly Lys Phe
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Val Trp Leu Lys Gly Glu Arg Lys Lys Arg Asp Arg Asp Glu Lys Asn
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tcccaaataa agcaaaacct agatccgaca ttgaaggaaa aaccttttag atccatctct 180
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Met Lys Arg Asp His His His His His Gln Asp Lys
1 5 10
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Lys Thr Met Met Met Asn Glu Glu Asp Asp Gly Asn Gly Met Asp Glu
15 20 25
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Leu Leu Ala Val Leu Gly Tyr Lys Val Arg Ser Ser Glu Met Ala Asp
30 35 40
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45 50 55 60
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Glu Asp Asp Leu Ser Gln Leu Ala Thr Glu Thr Val His Tyr Asn Pro
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mbil9 Sequence Listing.ST25															
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Pro	Ser	Ser	Asn	Ala	Glu	Tyr	Asp	Leu	Lys	Ala	Ile	Pro	Gly	Asp	Ala
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att	ctc	aat	cag	ttc	gct	atc	gat	tcg	gct	tct	tcg	tct	aac	caa	ggc
Ile	Leu	Asn	Gln	Phe	Ala	Ile	Asp	Ser	Ala	Ser	Ser	Ser	Asn	Gln	Gly
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Gly	Gly	Gly	Asp	Thr	Tyr	Thr	Thr	Asn	Lys	Arg	Leu	Lys	Cys	Ser	Asn
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Gly	Val	Val	Glu	Thr	Thr	Thr	Ala	Thr	Ala	Glu	Ser	Thr	Arg	His	Val
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Val	Leu	Val	Asp	Ser	Gln	Glu	Asn	Gly	Val	Arg	Leu	Val	His	Ala	Leu
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Ala	Leu	Val	Lys	Gln	Ile	Gly	Phe	Leu	Ala	Val	Ser	Gln	Ile	Gly	Ala
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tac	cgt	ctc	tct	ccg	tcg	cag	agt	cca	atc	gac	cac	tct	ctc	tcc	gat
Tyr	Arg	Leu	Ser	Pro	Ser	Gln	Ser	Pro	Ile	Asp	His	Ser	Leu	Ser	Asp
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Thr	Leu	Gln	Met	His	Phe	Tyr	Glu	Thr	Cys	Pro	Tyr	Leu	Lys	Phe	Ala
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His	Phe	Thr	Ala	Asn	Gln	Ala	Ile	Leu	Glu	Ala	Phe	Gln	Gly	Lys	Lys
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Arg	Val	His	Val	Ile	Asp	Phe	Ser	Met	Ser	Gln	Gly	Leu	Gln	Trp	Pro
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Arg	Leu	Thr	Gly	Ile	Gly	Pro	Pro	Ala	Pro	Asp	Asn	Phe	Asp	Tyr	Leu
				305					310					315	
cat	gaa	gtt	ggg	tgt	aag	ctg	gct	cat	tta	gct	gag	gcg	att	cac	gtt
His	Glu	Val	Gly	Cys	Lys	Leu	Ala	His	Leu	Ala	Glu	Ala	Ile	His	Val
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Glu	Phe	Glu	Tyr	Arg	Gly	Phe	Val	Ala	Asn	Thr	Leu	Ala	Asp	Leu	Asp
		335					340					345			
gct	tcg	atg	ctt	gag	ctt	aga	cca	agt	gag	att	gaa	tct	gtt	gcg	gtt
Ala	Ser	Met	Leu	Glu	Leu	Arg	Pro	Ser	Glu	Ile	Glu	Ser	Val	Ala	Val
	350					355					360				
aac	tct	gtt	ttc	gag	ctt	cac	aag	ctc	ttg	gga	cga	cct	ggt	gcg	atc
Asn	Ser	Val	Phe	Glu	His	Lys	Leu	Leu	Gly	Arg	Pro	Gly	Ala	Ile	
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 Val Val Glu Gln Glu Ser Asn His Asn Ser Pro Ile Phe Leu Asp Arg 410
 400 405

ttt act gag tcg ttg cat tat tac tcg acg ttg ttt gac tcg ttg gaa 1479
 Phe Thr Glu Ser Leu His Tyr Tyr Ser Thr Leu Phe Asp Ser Leu Glu 425
 415 420

ggg gta ccg agt ggt caa gac aag gtc atg tcg gag gtt tac ttg ggt 1527
 Gly Val Pro Ser Gly Gln Asp Lys Val Met Ser Glu Val Tyr Leu Gly 440
 430 435

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 Lys Gln Ile Cys Asn Val Val Ala Cys Asp Gly Pro Asp Arg Val Glu 460
 445 450 455

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 Arg His Glu Thr Leu Ser Gln Trp Arg Asn Arg Phe Gly Ser Ala Gly 475
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 495 500

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 510 515

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 Ser Ala Trp Lys Leu Ser Thr Asn 530
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tctgttgaac cggttatgat gatagatttc cgaccgaagc caaactaaat cctactgttt 1874

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Leu Glu Gln Leu Glu Val Met Met Ser Asn Val Gln Glu Asp Asp Leu
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Ser Gln Leu Ala Thr Glu Thr Val His Tyr Asn Pro Ala Glu Leu Tyr
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Thr Trp Leu Asp Ser Met Leu Thr Asp Leu Asn Pro Pro Ser Ser Asn

85 90 95

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 420 425 430
 Gly Gln Asp Lys Val Met Ser Glu Val Tyr Leu Gly Lys Gln Ile Cys
 435 440 445
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 450 455 460
 Leu Ser Gln Trp Arg Asn Arg Phe Gly Ser Ala Gly Phe Ala Ala Ala
 465 470 475 480
 His Ile Gly Ser Asn Ala Phe Lys Gln Ala Ser Met Leu Leu Ala Leu
 485 490 495
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 Thr Val Asp Val Glu Glu Leu Ser Val Glu Glu Arg Asn Leu Leu Ser
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 Val Ala Tyr Lys Asn Val Ile Gly Ala Arg Arg Ala Ser Trp Arg Ile
 50 55 60
 att tct tcg att gag cag aaa gaa gag agc aaa ggg aac gaa gat cat 240
 Ile Ser Ser Ile Glu Gln Lys Glu Glu Ser Lys Gly Asn Glu Asp His
 65 70 75 80
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mbil9 Sequence Listing.ST25

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tct gct tca cca gct gaa tct aaa gtg ttt tat ctt aag atg aag ggt      384
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210                      215                      220

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225                      230                      235                      240

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Ile Ser Ser Ile Glu Gln Lys Glu Glu Ser Lys Gly Asn Glu Asp His
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Val Ala Ile Ile Lys Asp Tyr Arg Gly Glu Ile Glu Ser Glu Leu Ser
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mbil9 Sequence Listing.ST25

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Ala Cys Ser Leu Ala Lys Gln Ala Phe Asp Asp Ala Ile Ala Glu Leu
 195 200 205

Asp Thr Leu Gly Glu Glu Ser Tyr Lys Asp Ser Thr Leu Ile Met Gln
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 25 30 35
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 Leu Arg Leu Tyr Gly Arg Ala Trp Gln Arg Ile Glu Glu His Ile Gly
 40 45 50

mbi19 Sequence Listing.ST25

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mbi19 Sequence Listing.ST25

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ggc cgt ata gcg ttt cag gct ctc ttt gca aga gaa aga ttg cct caa Gly Arg Ile Ala Phe Gln Ala Leu Phe Ala Arg Glu Arg Leu Pro Gln 535 540 545 550			1987
agc ttt tcg cct cct caa gtg gca gag aat gtg aat aga aaa caa agt Ser Phe Ser Pro Pro Gln Val Ala Glu Asn Val Asn Arg Lys Gln Ser 555 560 565			2035
gac acg tca atg cca ttg gct cct aat ttc aaa agc cag gat tct tgt Asp Thr Ser Met Pro Leu Ala Pro Asn Phe Lys Ser Gln Asp Ser Cys 570 575 580			2083
gct gca gac caa gaa gga gta gta atg atc ggt gtt gga aca tgc aag Ala Ala Asp Gln Glu Gly Val Val Met Ile Gly Val Gly Thr Cys Lys 585 590 595			2131
agt ctt aaa acg aga cag aca gga ttt aag cca tac aag aga tgt tca Ser Leu Lys Thr Arg Gln Thr Gly Phe Lys Pro Tyr Lys Arg Cys Ser 600 605 610			2179
atg gaa gtg aaa gag agc caa gtt ggg aac ata aac aat caa agt gat Met Glu Val Lys Glu Ser Gln Val Gly Asn Ile Asn Asn Gln Ser Asp 615 620 625 630			2227
gaa aaa gtc tgc aaa agg ctt cga ttg gaa gga gaa gct tct aca tga Glu Lys Val Cys Lys Arg Leu Arg Leu Glu Gly Glu Ala Ser Thr 635 640 645			2275
cagacttgga ggtaaaaaaa aaacatccac atttttatca atatctttaa atctagtggt			2335

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agtagtttgc ttctccaatc tttatgaaag agacttttaa ttttccttcc gaacatttct 2395
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 35 40 45

Ile Glu Glu His Ile Gly Thr Lys Thr Ala Val Gln Ile Arg Ser His
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Lys Arg Lys Pro Asn Thr Pro Tyr Pro Arg Lys Pro Gly Asn Asn Gly
 100 105 110

Thr Ser Ser Ser Gln Val Ser Ser Ala Lys Asp Ala Lys Leu Val Ser
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 130 135 140

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Gln Val Ser Gly Asp Ile Glu Thr Ser Lys Thr Ser Thr Val Asp Asn
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Ala Val Gln Asp Val Pro Lys Lys Asn Lys Asp Lys Asp Gly Asn Asp
 195 200 205

Gly Thr Thr Val His Ser Met Gln Asn Tyr Pro Trp His Phe His Ala
 210 215 220

Asp Ile Val Asn Gly Asn Ile Ala Lys Cys Pro Gln Asn His Pro Ser
 225 230 235 240

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Ser	Phe	Leu	Gln	Ile	Ser	Ser	Thr	Phe	Ser	Asn	Leu	Ile	Met	Ser	Thr
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Leu	Leu	Gln	Asn	Pro	Ala	Ala	His	Ala	Ala	Ala	Thr	Phe	Ala	Ala	Ser
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Val	Trp	Pro	Tyr	Ala	Ser	Val	Gly	Asn	Ser	Gly	Asp	Ser	Ser	Thr	Pro
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Met	Ser	Ser	Ser	Pro	Pro	Ser	Ile	Thr	Ala	Ile	Ala	Ala	Ala	Thr	Val
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Glu	Lys	Gln	Asn	Thr	Ala	Leu	Gln	Asp	Gln	Thr	Leu	Ala	Ser	Lys	Ser
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Pro	Ala	Ser	Ser	Ser	Asp	Asp	Ser	Asp	Glu	Thr	Gly	Val	Thr	Lys	Leu
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Asn	Ala	Asp	Ser	Lys	Thr	Asn	Asp	Asp	Lys	Ile	Glu	Glu	Val	Val	Val
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Thr	Ala	Ala	Val	His	Asp	Ser	Asn	Thr	Ala	Gln	Lys	Lys	Asn	Leu	Val
	450					455					460				
Asp	Arg	Ser	Ser	Cys	Gly	Ser	Asn	Thr	Pro	Ser	Gly	Ser	Asp	Ala	Glu
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Thr	Asp	Ala	Leu	Asp	Lys	Met	Glu	Lys	Asp	Lys	Glu	Asp	Val	Lys	Glu
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Thr	Asp	Glu	Asn	Gln	Pro	Asp	Val	Ile	Glu	Leu	Asn	Asn	Arg	Lys	Ile
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Lys	Met	Arg	Asp	Asn	Asn	Ser	Asn	Asn	Asn	Ala	Thr	Thr	Asp	Ser	Trp
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Lys	Glu	Val	Ser	Glu	Glu	Gly	Arg	Ile	Ala	Phe	Gln	Ala	Leu	Phe	Ala

540

mbi19 Sequence Listing.ST25

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Thr Tyr Asn Glu Glu Leu Glu Gln Ser Lys Arg Arg Arg Asn Gly Asn			
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Gly Asn Met Thr Arg Thr Leu Leu Thr Ser Gly Leu Ser Asn Asp Gly			
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gtt tct acg acg ggg ttt aga tcg gcg gag gca ctg ttt gag aaa gcg			636
Val Ser Thr Thr Gly Phe Arg Ser Ala Glu Ala Leu Phe Glu Lys Ala			
180	185	190	
gta acg cca agc gac gtt ggg aag cta aac cgt ttg gtt ata ccg aaa			684
Val Thr Pro Ser Asp Val Gly Lys Leu Asn Arg Leu Val Ile Pro Lys			
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His His Ala Glu Lys His Phe Pro Leu Pro Ser Ser Asn Val Ser Val			
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Lys Gly Val Leu Leu Asn Phe Glu Asp Val Asn Gly Lys Val Trp Arg			
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Phe Arg Tyr Ser Tyr Trp Asn Ser Ser Gln Ser Tyr Val Leu Thr Lys			
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Val Ser Phe Ser Arg Ser Asn Gly Gln Asp Gln Gln Leu Tyr Ile Gly			
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290	295	300	
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Leu Phe Gly Val Asn Ile Ser Pro Glu Ser Ser Arg Asn Asp Val Val			
305	310	315	
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Gly Asn Lys Arg Val Asn Asp Thr Glu Met Leu Ser Leu Val Cys Ser			
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Lys Lys Gln Arg Ile Phe His Ala Ser			
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ttctttcttc ttgtttacca aaggttcatg agttgttttt gttgtattga tgaactgtaa			1238
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<213> Arabidopsis thaliana

<400> 22

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mbi19 Sequence Listing.ST25

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 His Gln Arg Val Trp Leu Gly Thr Phe Asn Glu Glu Asp Glu Ala Ala
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 Arg Ala Tyr Asp Val Ala Val His Arg Phe Arg Arg Arg Asp Ala Val
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 Thr Asn Phe Lys Asp Val Lys Met Asp Glu Asp Glu Val Asp Phe Leu
 115 120 125
 Asn Ser His Ser Lys Ser Glu Ile Val Asp Met Leu Arg Lys His Thr
 130 135 140
 Tyr Asn Glu Glu Leu Glu Gln Ser Lys Arg Arg Arg Asn Gly Asn Gly
 145 150 155 160
 Asn Met Thr Arg Thr Leu Leu Thr Ser Gly Leu Ser Asn Asp Gly Val
 165 170 175
 Ser Thr Thr Gly Phe Arg Ser Ala Glu Ala Leu Phe Glu Lys Ala Val
 180 185 190
 Thr Pro Ser Asp Val Gly Lys Leu Asn Arg Leu Val Ile Pro Lys His
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 His Ala Glu Lys His Phe Pro Leu Pro Ser Ser Asn Val Ser Val Lys
 210 215 220
 Gly Val Leu Leu Asn Phe Glu Asp Val Asn Gly Lys Val Trp Arg Phe
 225 230 235 240
 Arg Tyr Ser Tyr Trp Asn Ser Ser Gln Ser Tyr Val Leu Thr Lys Gly
 245 250 255
 Trp Ser Arg Phe Val Lys Glu Lys Asn Leu Arg Ala Gly Asp Val Val
 260 265 270
 Ser Phe Ser Arg Ser Asn Gly Gln Asp Gln Gln Leu Tyr Ile Gly Trp
 275 280 285
 Lys Ser Arg Ser Gly Ser Asp Leu Asp Ala Gly Arg Val Leu Arg Leu
 290 295 300
 Phe Gly Val Asn Ile Ser Pro Glu Ser Ser Arg Asn Asp Val Val Gly
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mbi19 Sequence Listing.ST25

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724

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Thr Arg His Pro Ile Tyr Arg Gly Val Arg Gln Arg Asn Ser Gly Lys
 50 55 60

Trp Val Cys Glu Val Arg Glu Pro Asn Lys Lys Ser Arg Ile Trp Leu
 65 70 75 80

Gly Thr Phe Pro Thr Val Glu Met Ala Ala Arg Ala His Asp Val Ala
 85 90 95

Ala Leu Ala Leu Arg Gly Arg Ser Ala Cys Leu Asn Phe Ala Asp Ser
 100 105 110

Ala Trp Arg Leu Arg Ile Pro Glu Thr Thr Cys Pro Lys Glu Ile Gln
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Lys Ala Ala Ser Glu Ala Ala Met Ala Phe Gln Asn Glu Thr Thr Thr
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Glu Gly Ser Lys Thr Ala Ala Glu Ala Glu Glu Ala Ala Gly Glu Gly
 145 150 155 160

Val Arg Glu Gly Glu Arg Arg Ala Glu Glu Gln Asn Gly Gly Val Phe
 165 170 175

Tyr Met Asp Asp Glu Ala Leu Leu Gly Met Pro Asn Phe Phe Glu Asn
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Met Ala Glu Gly Met Leu Leu Pro Pro Pro Glu Val Gly Trp Asn His
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mbi19 Sequence Listing.ST25

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 275 280 285 290

aca ata tcg aat cat tca aat tcg tcc tta tac agt gat ata aaa tca 970
 Thr Ile Ser Asn His Ser Asn Ser Ser Leu Tyr Ser Asp Ile Lys Ser
 295 300 305

gag acc aat ttt ttt ggc aca gag gct aca aat gtt ggt atg tgg cca 1018
 Glu Thr Asn Phe Phe Gly Thr Glu Ala Thr Asn Val Gly Met Trp Pro
 310 315 320

tgt aac cag ctt cag cct cag caa cat gca tat ggc cat ata taa 1063
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 35 40 45

Cys Gly Lys Ser Cys Arg Leu Arg Trp Ile Asn Tyr Leu Arg Pro Asp
 50 55 60

Leu Lys Arg Gly Ala Phe Ser Gln Asp Glu Glu Asn Leu Ile Ile Glu
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Leu His Ala Val Leu Gly Asn Arg Trp Ser Gln Ile Ala Ala Gln Leu
 85 90 95

Pro Gly Arg Thr Asp Asn Glu Ile Lys Asn Leu Trp Asn Ser Cys Leu
 100 105 110

Lys Lys Lys Leu Arg Leu Arg Gly Ile Asp Pro Val Thr His Lys Leu
 115 120 125

Leu Thr Glu Ile Glu Thr Gly Thr Asp Asp Lys Thr Lys Pro Val Glu
 130 135 140

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 145 150 155 160

Thr Thr Cys Ser Thr Asn Gln Asn Asn Asn Thr Asp His Leu Tyr Thr
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mbi19 Sequence Listing.ST25

Gly Asn Phe Gly Phe Gln Arg Leu Ser Leu Glu Asn Gly Ser Arg Ile
 180 185 190

Ala Ala Gly Ser Asp Leu Gly Ile Trp Ile Pro Gln Thr Gly Arg Asn
 195 200 205

His His His His Val Asp Glu Thr Ile Pro Ser Ala Val Val Leu Pro
 210 215 220

Gly Ser Met Phe Ser Ser Gly Leu Thr Gly Tyr Arg Ser Ser Asn Leu
 225 230 235 240

Gly Leu Ile Glu Leu Glu Asn Ser Phe Ser Thr Gly Pro Met Met Thr
 245 250 255

Glu His Gln Gln Ile Gln Glu Ser Asn Tyr Asn Asn Ser Thr Phe Phe
 260 265 270

Gly Asn Gly Asn Leu Asn Trp Gly Leu Thr Met Glu Glu Asn Gln Asn
 275 280 285

Pro Phe Thr Ile Ser Asn His Ser Asn Ser Ser Leu Tyr Ser Asp Ile
 290 295 300

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 Gln His Leu Gln Gln Gln Gln Gln Pro Pro Pro Gly Met Leu Met Ser
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 His His Asn Ser Tyr Asn Arg Asn Pro Asn Ala Ala Ala Ala Val Leu
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 atg ggt cac aac acc tcc aca tct caa gct atg cat caa aga tta cct 365
 Met Gly His Asn Thr Ser Thr Ser Gln Ala Met His Gln Arg Leu Pro
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mbi19 Sequence Listing.ST25

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mbil9 Sequence Listing.ST25

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 Pro Asn Ala Ala Ala Ala Val Leu Met Gly His Asn Thr Ser Thr Ser
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 Gln Ala Met His Gln Arg Leu Pro Phe Gly Gly Ser Met Ser Pro His
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 Asp Gln Lys Thr Leu Glu Ser Leu Gly Phe Pro Thr Ser Pro Leu Pro
 100 105 110
 Ser Ala Ser Asn Ser Tyr Gly Gly Gly Asn Glu Gly Gly Gly Gly
 115 120 125
 Asp Ser Ala Gly Ala Asn Ala Asn Ser Ser Asp Pro Pro Ala Lys Arg
 130 135 140
 Asn Arg Gly Arg Pro Pro Gly Ser Gly Lys Lys Gln Leu Asp Ala Leu
 145 150 155 160
 Gly Gly Thr Gly Gly Val Gly Phe Thr Pro His Val Ile Glu Val Lys
 165 170 175
 Thr Gly Glu Asp Ile Ala Thr Lys Ile Leu Ala Phe Thr Asn Gln Gly
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 Pro Arg Ala Ile Cys Ile Leu Ser Ala Thr Gly Ala Val Thr Asn Val
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mbil9 Sequence Listing.ST25

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Glu Ser Asn Gly Thr Val Thr Lys Thr Gly Asn Leu Ser Val Ser Leu
245 250 255

Ala Gly His Glu Gly Arg Ile Val Gly Gly Cys Val Asp Gly Met Leu
260 265 270

Val Ala Gly Ser Gln Val Gln Val Ile Val Gly Ser Phe Val Pro Asp
275 280 285

Gly Arg Lys Gln Lys Gln Ser Ala Gly Arg Ala Gln Asn Thr Pro Glu
290 295 300

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325 330 335

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Met Ser Ser Ser Glu Arg
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25 30 35

mbi19 Sequence Listing.ST25																
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cga Arg	gtg Val 280	gaa Glu	gct Ala	gca Ala	tat Tyr	gta Val 285	ggg Gly	aaa Lys	ggg Gly	gct Ala	gct Ala 290	tct Ser	tca Ser	ttc Phe	aca Thr	978
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ggc Gly	cag Gln	gta Val	caa Gln 330	cca Pro	aca Thr	aaa Lys	tct Ser	gag Glu 335	agc Ser	aac Asn	aat Asn	cgt Arg	cca Pro 340	att Ile	acc Thr	1122

mbi19 Sequence Listing.ST25

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Glu His Ile Ala Gly Thr Ser Cys Lys Thr Thr Arg Leu Val Ala Thr
      360                      365                      370

aag gct gat ctg gag cgg ctg gct cag aac aga gga gat gca atg cag      1266
Lys Ala Asp Leu Glu Arg Leu Ala Gln Asn Arg Gly Asp Ala Met Gln
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cgt tac aag gaa aag agg aag aca cgg aga tat gat aag acc ata agg      1314
Arg Tyr Lys Glu Lys Arg Lys Thr Arg Arg Tyr Asp Lys Thr Ile Arg
      395                      400                      405

tat gaa tcg agg aag gca aga gct gac act agg ttg cgt gtc aga ggc      1362
Tyr Glu Ser Arg Lys Ala Arg Ala Asp Thr Arg Leu Arg Val Arg Gly
      410                      415                      420

aga ttt gtg aaa gct agt gaa gct cct tac cct taa ccttaagttt      1408
Arg Phe Val Lys Ala Ser Glu Ala Pro Tyr Pro
      425                      430

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35      40      45

Ser Gln Ile Cys Asp Asn Cys Gly Asn Glu Pro Val Ser Val Arg Cys
50      55      60

Phe Thr Asp Asn Leu Ile Leu Cys Gln Glu Cys Asp Trp Asp Val His
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Gly Ser Cys Ser Val Ser Asp Ala His Val Arg Ser Ala Val Glu Gly
85      90      95

Phe Ser Gly Cys Pro Ser Ala Leu Glu Leu Ala Ala Leu Trp Gly Leu
100     105     110

Asp Leu Glu Gln Gly Arg Lys Asp Glu Glu Asn Gln Val Pro Met Met
115     120     125

Ala Met Met Met Asp Asn Phe Gly Met Gln Leu Asp Ser Trp Val Leu
130     135     140

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mbi19 Sequence Listing.ST25

Gly Ser Asn Glu Leu Ile Val Pro Ser Asp Thr Thr Phe Lys Lys Arg
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 165 170 175

Gln Leu Glu Glu Leu Leu Lys Ser Gly Val Val Gly Gly Asp Gly Asp
 180 185 190

Asp Gly Asp Arg Asp Arg Asp Cys Asp Arg Glu Gly Ala Cys Asp Gly
 195 200 205

Asp Gly Asp Gly Glu Ala Gly Glu Gly Leu Met Val Pro Glu Met Ser
 210 215 220

Glu Arg Leu Lys Trp Ser Arg Asp Val Glu Glu Ile Asn Gly Gly Gly
 225 230 235 240

Gly Gly Gly Val Asn Gln Gln Trp Asn Ala Thr Thr Thr Asn Pro Ser
 245 250 255

Gly Gly Gln Ser Ser Gln Ile Trp Asp Phe Asn Leu Gly Gln Ser Arg
 260 265 270

Gly Pro Glu Asp Thr Ser Arg Val Glu Ala Ala Tyr Val Gly Lys Gly
 275 280 285

Ala Ala Ser Ser Phe Thr Ile Asn Asn Phe Val Asp His Met Asn Glu
 290 295 300

Thr Cys Ser Thr Asn Val Lys Gly Val Lys Glu Ile Lys Lys Asp Asp
 305 310 315 320

Tyr Lys Arg Ser Thr Ser Gly Gln Val Gln Pro Thr Lys Ser Glu Ser
 325 330 335

Asn Asn Arg Pro Ile Thr Phe Gly Ser Glu Lys Gly Ser Asn Ser Ser
 340 345 350

Ser Asp Leu His Phe Thr Glu His Ile Ala Gly Thr Ser Cys Lys Thr
 355 360 365

Thr Arg Leu Val Ala Thr Lys Ala Asp Leu Glu Arg Leu Ala Gln Asn
 370 375 380

Arg Gly Asp Ala Met Gln Arg Tyr Lys Glu Lys Arg Lys Thr Arg Arg
 385 390 395 400

Tyr Asp Lys Thr Ile Arg Tyr Glu Ser Arg Lys Ala Arg Ala Asp Thr
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mbi19 Sequence Listing.ST25

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gag gag ata cag caa cag caa cag gaa cag caa cag caa cag ctg caa Glu Glu Ile Gln Gln Gln Gln Glu Gln Gln Gln Gln Gln Leu Gln 235 240 245			954
ccg gat ttg ctt act gtt gca gat tac ggt tgg cct tgg tct aat gat Pro Asp Leu Leu Thr Val Ala Asp Tyr Gly Trp Pro Trp Ser Asn Asp 250 255 260 265			1002
att gta aat gat cag act tct tgg gat cct aat gag tgc ttt gat att Ile Val Asn Asp Gln Thr Ser Trp Asp Pro Asn Glu Cys Phe Asp Ile 270 275 280			1050
aat gaa ctc ctt gga gat ttg aat gaa cct ggt ccc cat cag agc caa Asn Glu Leu Leu Gly Asp Leu Asn Glu Pro Gly Pro His Gln Ser Gln 285 290 295			1098
gac caa aac cac gta aat tct ggt agt tat gat ttg cat ccg ctt cat Asp Gln Asn His Val Asn Ser Gly Ser Tyr Asp Leu His Pro Leu His 300 305 310			1146
ctc gag cca cac gat ggt cac gag ttc aat ggt ttg agt tct ctg gat Leu Glu Pro His Asp Gly His Glu Phe Asn Gly Leu Ser Ser Leu Asp 315 320 325			1194
att tga gagttctgag gcaatggtcc tacaagacta caacataatc tttggattga Ile 330			1250
tcataggaga aacaagaaat aggtgttaat gatctgattc acaatgaaaa aatatttaat			1310
aactctatag tttttgttct ttccttgat catgaactgt tgcttctcat ctattgagtt			1370
aatatagcga atagcagagt ttctctcata aaaaaaaaaa aaa			1413
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Trp Lys Glu Tyr Asn Glu Ile Val Glu Ala Ser Ala Val Lys Glu Gly 35 40 45			
Glu Lys Pro Lys Arg Lys Val Pro Ala Lys Gly Ser Lys Lys Gly Cys 50 55 60			
Met Lys Gly Lys Gly Gly Pro Asp Asn Ser His Cys Ser Phe Arg Gly 65 70 75 80			
Val Arg Gln Arg Ile Trp Gly Lys Trp Val Ala Glu Ile Arg Glu Pro 85 90 95			
Lys Ile Gly Thr Arg Leu Trp Leu Gly Thr Phe Pro Thr Ala Glu Lys 100 105 110			

mbi19 Sequence Listing.ST25

Ala Ala Ser Ala Tyr Asp Glu Ala Ala Thr Ala Met Tyr Gly Ser Leu
 115 120 125

Ala Arg Leu Asn Phe Pro Gln Ser Val Gly Ser Glu Phe Thr Ser Thr
 130 135 140

Ser Ser Gln Ser Glu Val Cys Thr Val Glu Asn Lys Ala Val Val Cys
 145 150 155 160

Gly Asp Val Cys Val Lys His Glu Asp Thr Asp Cys Glu Ser Asn Pro
 165 170 175

Phe Ser Gln Ile Leu Asp Val Arg Glu Glu Ser Cys Gly Thr Arg Pro
 180 185 190

Asp Ser Cys Thr Val Gly His Gln Asp Met Asn Ser Ser Leu Asn Tyr
 195 200 205

Asp Leu Leu Leu Glu Phe Glu Gln Gln Tyr Trp Gly Gln Val Leu Gln
 210 215 220

Glu Lys Glu Lys Pro Lys Gln Glu Glu Glu Glu Ile Gln Gln Gln Gln
 225 230 235 240

Gln Glu Gln Gln Gln Gln Gln Leu Gln Pro Asp Leu Leu Thr Val Ala
 245 250 255

Asp Tyr Gly Trp Pro Trp Ser Asn Asp Ile Val Asn Asp Gln Thr Ser
 260 265 270

Trp Asp Pro Asn Glu Cys Phe Asp Ile Asn Glu Leu Leu Gly Asp Leu
 275 280 285

Asn Glu Pro Gly Pro His Gln Ser Gln Asp Gln Asn His Val Asn Ser
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Gly Ser Tyr Asp Leu His Pro Leu His Leu Glu Pro His Asp Gly His
 305 310 315 320

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 Ala Ile Gln Ser His Leu Leu Glu Asp Leu Leu Val Cys Asp Gly Phe

mbi19 Sequence Listing.ST25

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30	35	40	
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Ile Glu Pro His Val Pro Lys Gln Glu Pro Asp Ser Pro Val Leu Asp			
45	50	55	
ccg gat tct ttc gtc aac gag ttc ttg caa gtg gaa ggg gaa tca tca			242
Pro Asp Ser Phe Val Asn Glu Phe Leu Gln Val Glu Gly Glu Ser Ser			
60	65	70	75
tca tca tca tca cca gag ctg aat tca tct tca tca aca tat gag act			290
Ser Ser Ser Ser Pro Glu Leu Asn Ser Ser Ser Ser Thr Tyr Glu Thr			
80	85	90	
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Asp Gln Ser Val Lys Lys Ala Glu Arg Phe Glu Glu Glu Val Asp Ala			
95	100	105	
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Arg His Tyr Arg Gly Val Arg Arg Arg Pro Trp Gly Lys Phe Ala Ala			
110	115	120	
gag att cga gat cca gca aag aaa gga tca aga atc tgg cta gga aca			434
Glu Ile Arg Asp Pro Ala Lys Lys Gly Ser Arg Ile Trp Leu Gly Thr			
125	130	135	
ttt gag agt gat gtt gat gct gca aga gcc tat gac tgt gca gct ttc			482
Phe Glu Ser Asp Val Asp Ala Ala Arg Ala Tyr Asp Cys Ala Ala Phe			
140	145	150	155
aag ctc cgg gga aga aaa gcc gtg ctc aac ttc cct ctt gac gcc ggg			530
Lys Leu Arg Gly Arg Lys Ala Val Leu Asn Phe Pro Leu Asp Ala Gly			
160	165	170	
aaa tat gaa gct cca gcg aat tca gga agg aaa agg aag aga agt gat			578
Lys Tyr Glu Ala Pro Ala Asn Ser Gly Arg Lys Arg Lys Arg Ser Asp			
175	180	185	
gtg cat gaa gag ctt caa aga act cag agc aat tca tct tca tct tcc			626
Val His Glu Glu Leu Gln Arg Thr Gln Ser Asn Ser Ser Ser Ser Ser			
190	195	200	
tgt gat gca ttt tag catattaaga gtgtgagcag tttccttaag ttgtataaag			681
Cys Asp Ala Phe			
205			
taattgtaca gaggaacga attgtgtagg tttagtgtgc ttgcaagttg caacaaatgt			741
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20	25	30	
Phe Asp Ala Ser Phe Val Ser Gly Leu Trp Cys Ile Glu Pro His Val			
35	40	45	

mbil9 Sequence Listing.ST25

Pro Lys Gln Glu Pro Asp Ser Pro Val Leu Asp Pro Asp Ser Phe Val
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Asn Glu Phe Leu Gln Val Glu Gly Glu Ser Ser Ser Ser Ser Ser Pro
 65 70 75 80

Glu Leu Asn Ser Ser Ser Ser Thr Tyr Glu Thr Asp Gln Ser Val Lys
 85 90 95

Lys Ala Glu Arg Phe Glu Glu Glu Val Asp Ala Arg His Tyr Arg Gly
 100 105 110

Val Arg Arg Arg Pro Trp Gly Lys Phe Ala Ala Glu Ile Arg Asp Pro
 115 120 125

Ala Lys Lys Gly Ser Arg Ile Trp Leu Gly Thr Phe Glu Ser Asp Val
 130 135 140

Asp Ala Ala Arg Ala Tyr Asp Cys Ala Ala Phe Lys Leu Arg Gly Arg
 145 150 155 160

Lys Ala Val Leu Asn Phe Pro Leu Asp Ala Gly Lys Tyr Glu Ala Pro
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 Met Ala Asp Arg Ile Lys Gly Pro Trp Ser Pro Glu Glu Asp Glu
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cag ctt cgt agg ctt gtt gtt aaa tac ggt cca aga aac tgg aca gtg 158
 Gln Leu Arg Arg Leu Val Val Lys Tyr Gly Pro Arg Asn Trp Thr Val
 20 25 30

att agc aaa tct att ccc ggt aga tcg ggg aaa tcg tgt cgt tta cgg 206
 Ile Ser Lys Ser Ile Pro Gly Arg Ser Gly Lys Ser Cys Arg Leu Arg
 35 40 45

tgg tgc aac cag ctt tcg ccg caa gtt gag cat cgg ccg ttt tcg gct 254
 Trp Cys Asn Gln Leu Ser Pro Gln Val Glu His Arg Pro Phe Ser Ala
 50 55 60

gag gaa gac gag acg atc gca cgt gct cac gct cag ttc ggg aat aaa 302
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 65 70 75

mbil9 Sequence Listing.ST25

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80 85 90 95	
aag aat cac tgg aac tcg acg ctc aag agg aaa tgc ggc ggt tac gac	398
Lys Asn His Trp Asn Ser Thr Leu Lys Arg Lys Cys Gly Gly Tyr Asp	
100 105 110	
cat cgg ggt tac gat ggt tcg gag gat cat cgg ccg gtt aag aga tcg	446
His Arg Gly Tyr Asp Gly Ser Glu Asp His Arg Pro Val Lys Arg Ser	
115 120 125	
gtg agt gcg gga tct cca cct gtt gtt act ggg ctt tac atg agc cca	494
Val Ser Ala Gly Ser Pro Pro Val Val Thr Gly Leu Tyr Met Ser Pro	
130 135 140	
gga agc cca act gga tct gat gtc agt gat tca agt act atc ccg ata	542
Gly Ser Pro Thr Gly Ser Asp Val Ser Asp Ser Ser Thr Ile Pro Ile	
145 150 155	
tta cct tcc gtt gag ctt ttc aag cct gtg cct aga cct ggt gct gtt	590
Leu Pro Ser Val Glu Leu Phe Lys Pro Val Pro Arg Pro Gly Ala Val	
160 165 170 175	
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Val Leu Pro Leu Pro Ile Glu Thr Ser Ser Phe Ser Asp Asp Pro Pro	
180 185 190	
act tcg tta agc ttg tca ctt cct ggt gcc gac gta agc gag gag tca	686
Thr Ser Leu Ser Leu Ser Leu Pro Gly Ala Asp Val Ser Glu Glu Ser	
195 200 205	
aac cgt agc cac gag tca acg aat atc aac aac acc act tcg agc cgc	734
Asn Arg Ser His Glu Ser Thr Asn Ile Asn Asn Thr Ser Ser Arg	
210 215 220	
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His Asn His Asn Asn Thr Ser Phe Met Pro Phe Ser Gly Gly Phe	
225 230 235	
aga ggt gcg att gag gaa atg ggg aag tct ttt ccc ggt aac gga ggc	830
Arg Gly Ala Ile Glu Glu Met Gly Lys Ser Phe Pro Gly Asn Gly Gly	
240 245 250 255	
gag ttt atg gcg gtg gtg caa gag atg att aag gcg gaa gtg agg agt	878
Glu Phe Met Ala Val Val Gln Glu Met Ile Lys Ala Glu Val Arg Ser	
260 265 270	
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Tyr Met Thr Glu Met Gln Arg Asn Asn Gly Gly Gly Phe Val Gly Gly	
275 280 285	
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Phe Ile Asp Asn Gly Met Ile Pro Met Ser Gln Ile Gly Val Gly Arg	
290 295 300	
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Ile Glu	
305	
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aaatgtatag aggaaatcga gtgaacaaag ctcgagagct ggggacgtag tgacgaagac	1143
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 <212> PRT
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<400> 36

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mbi19 Sequence Listing.ST25

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	35	40	45
Cys Asn Gln	Leu Ser Pro	Gln Val Glu	His Arg Pro Phe Ser Ala Glu
	50	55	60
Glu Asp Glu	Thr Ile Ala	Arg Ala His	Ala Gln Phe Gly Asn Lys Trp
	65	70	75
Ala Thr Ile	Ala Arg	Leu Leu Asn	Gly Arg Thr Asp Asn Ala Val Lys
	85	90	95
Asn His Trp	Asn Ser Thr	Leu Lys Arg	Lys Cys Gly Gly Tyr Asp His
	100	105	110
Arg Gly Tyr	Asp Gly Ser	Glu Asp His	Arg Pro Val Lys Arg Ser Val
	115	120	125
Ser Ala Gly	Ser Pro Pro	Val Val Thr	Gly Leu Tyr Met Ser Pro Gly
	130	135	140
Ser Pro Thr	Gly Ser Asp	Val Ser Asp	Ser Ser Thr Ile Pro Ile Leu
	145	150	155
Pro Ser Val	Glu Leu Phe	Lys Pro Val	Pro Arg Pro Gly Ala Val Val
	165	170	175
Leu Pro Leu	Pro Ile Glu	Thr Ser Ser	Phe Ser Asp Asp Pro Pro Thr
	180	185	190
Ser Leu Ser	Leu Ser Leu	Pro Gly Ala	Asp Val Ser Glu Glu Ser Asn
	195	200	205
Arg Ser His	Glu Ser Thr	Asn Ile Asn	Asn Thr Thr Ser Ser Arg His
	210	215	220
Asn His Asn	Asn Thr Val	Ser Phe Met	Pro Phe Ser Gly Gly Phe Arg
	225	230	235
Gly Ala Ile	Glu Glu Met	Gly Lys Ser	Phe Pro Gly Asn Gly Gly Glu
	245	250	255
Phe Met Ala	Val Val Gln	Glu Met Ile	Lys Ala Glu Val Arg Ser Tyr
	260	265	270
Met Thr Glu	Met Gln Arg	Asn Asn Gly	Gly Gly Phe Val Gly Gly Phe
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	290	295	300

mbi19 Sequence Listing.ST25

Glu
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<223> G227

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Ile Lys Gly Pro Trp Ser Pro Glu Glu Asp Asp Leu Leu Gln Arg Leu
15 20 25

gtt cag aaa cat ggt ccg agg aac tgg tct ttg att agc aaa tca atc 149
Val Gln Lys His Gly Pro Arg Asn Trp Ser Leu Ile Ser Lys Ser Ile
30 35 40

cct gga cgt tcc ggc aaa tct tgt cgt ctc cgg tgg tgt aac cag cta 197
Pro Gly Arg Ser Gly Lys Ser Cys Arg Leu Arg Trp Cys Asn Gln Leu
45 50 55

tct ccg gag gta gag cac cgt gct ttt tcg cag gaa gaa gac gag acg 245
Ser Pro Glu Val Glu His Arg Ala Phe Ser Gln Glu Glu Asp Glu Thr
60 65 70 75

att att cga gct cac gct cgg ttt ggt aac aag tgg gct acg atc tct 293
Ile Ile Arg Ala His Ala Arg Phe Gly Asn Lys Trp Ala Thr Ile Ser
80 85 90

cgt ctt ctc aat gga cga acc gat aac gct atc aag aat cat tgg aac 341
Arg Leu Leu Asn Gly Arg Thr Asp Asn Ala Ile Lys Asn His Trp Asn
95 100 105

tcg acg ctg aag cga aaa tgc agc gtc gaa ggg caa agt tgt gat ttt 389
Ser Thr Leu Lys Arg Lys Cys Ser Val Glu Gly Gln Ser Cys Asp Phe
110 115 120

ggg ggt aat gga ggg tat gat ggt aat tta gga gaa gag caa ccg ttg 437
Gly Gly Asn Gly Gly Tyr Asp Gly Asn Leu Gly Glu Glu Gln Pro Leu
125 130 135

aaa cgt acg gcg agt ggt ggt ggt ggt gtc tcg act ggc ttg tat atg 485
Lys Arg Thr Ala Ser Gly Gly Gly Gly Val Ser Thr Gly Leu Tyr Met
140 145 150 155

agt ccc gga agt cca tcg gga tct gac gtc agc gag caa tct agt ggt 533
Ser Pro Gly Ser Pro Ser Gly Ser Asp Val Ser Glu Gln Ser Ser Gly
160 165 170

ggg gca cac gtg ttt aaa cca acg gtt aga tct gag gtt aca gcg tca 581
Gly Ala His Val Phe Lys Pro Thr Val Arg Ser Glu Val Thr Ala Ser
175 180 185

tcg tct ggt gaa gat cct cca act tat ctt agt ttg tct ctt cct tgg 629
Ser Ser Gly Glu Asp Pro Pro Thr Tyr Leu Ser Leu Ser Leu Pro Trp
190 195 200

act gac gag acg gtt cga gtc aac gag ccg gtt caa ctt aac cag aat 677
Thr Asp Glu Thr Val Arg Val Asn Glu Pro Val Gln Leu Asn Gln Asn
205 210 215

acg gtt atg gac ggt ggt tat acg gcg gag ctg ttt ccg gtt aga aag 725
Thr Val Met Asp Gly Gly Tyr Thr Ala Glu Leu Phe Pro Val Arg Lys
220 225 230 235

mbi19 Sequence Listing.ST25

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gaa gag caa gtg gaa gta gaa gaa gaa gaa gcg aag ggg ata tct ggt      773
Glu Glu Gln Val Glu Val Glu Glu Glu Glu Ala Lys Gly Ile Ser Gly
240\ 245 250

gga ttc ggt ggt gag ttc atg acg gtg gtt cag gag atg ata agg acg      821
Gly Phe Gly Gly Glu Phe Met Thr Val Val Gln Glu Met Ile Arg Thr
255 260 265

gag gtg agg agt tac atg gcg gat tta cag cga gga aac gtc ggt ggt      869
Glu Val Arg Ser Tyr Met Ala Asp Leu Gln Arg Gly Asn Val Gly Gly
270 275 280

agt agt tct ggc ggc gga ggt ggc ggt tgc tgt atg cca caa agt gta      917
Ser Ser Ser Gly Gly Gly Gly Gly Gly Ser Cys Met Pro Gln Ser Val
285 290 295

aac agc cgt cgt gtt ggg ttt aga gag ttt ata gtg aac caa atc gga      965
Asn Ser Arg Arg Val Gly Phe Arg Glu Phe Ile Val Asn Gln Ile Gly
300 305 310 315

att ggg aag atg gag tag gcggcc      989
Ile Gly Lys Met Glu
320

<210> 38
<211> 320
<212> PRT
<213> Arabidopsis thaliana

<400> 38

Met Ser Asn Pro Thr Arg Lys Asn Met Glu Arg Ile Lys Gly Pro Trp
1 5 10 15

Ser Pro Glu Glu Asp Asp Leu Leu Gln Arg Leu Val Gln Lys His Gly
20 25 30

Pro Arg Asn Trp Ser Leu Ile Ser Lys Ser Ile Pro Gly Arg Ser Gly
35 40 45

Lys Ser Cys Arg Leu Arg Trp Cys Asn Gln Leu Ser Pro Glu Val Glu
50 55 60

His Arg Ala Phe Ser Gln Glu Glu Asp Glu Thr Ile Ile Arg Ala His
65 70 75 80

Ala Arg Phe Gly Asn Lys Trp Ala Thr Ile Ser Arg Leu Leu Asn Gly
85 90 95

Arg Thr Asp Asn Ala Ile Lys Asn His Trp Asn Ser Thr Leu Lys Arg
100 105 110

Lys Cys Ser Val Glu Gly Gln Ser Cys Asp Phe Gly Gly Asn Gly Gly
115 120 125

Tyr Asp Gly Asn Leu Gly Glu Glu Gln Pro Leu Lys Arg Thr Ala Ser
130 135 140

Gly Gly Gly Gly Val Ser Thr Gly Leu Tyr Met Ser Pro Gly Ser Pro
145 150 155 160

Ser Gly Ser Asp Val Ser Glu Gln Ser Ser Gly Gly Ala His Val Phe
165 170 175

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mbil9 Sequence Listing.ST25

Lys Pro Thr Val Arg Ser Glu Val Thr Ala Ser Ser Ser Gly Glu Asp
180 185 190

Pro Pro Thr Tyr Leu Ser Leu Ser Leu Pro Trp Thr Asp Glu Thr Val
195 200 205

Arg Val Asn Glu Pro Val Gln Leu Asn Gln Asn Thr Val Met Asp Gly
210 215 220

Gly Tyr Thr Ala Glu Leu Phe Pro Val Arg Lys Glu Glu Gln Val Glu
225 230 235 240

Val Glu Glu Glu Glu Ala Lys Gly Ile Ser Gly Gly Phe Gly Gly Glu
245 250 255

Phe Met Thr Val Val Gln Glu Met Ile Arg Thr Glu Val Arg Ser Tyr
260 265 270

Met Ala Asp Leu Gln Arg Gly Asn Val Gly Gly Ser Ser Ser Gly Gly
275 280 285

Gly Gly Gly Gly Ser Cys Met Pro Gln Ser Val Asn Ser Arg Arg Val
290 295 300

Gly Phe Arg Glu Phe Ile Val Asn Gln Ile Gly Ile Gly Lys Met Glu
305 310 315 320

<210> 39
<211> 994
<212> DNA
<213> Arabidopsis thaliana

<220>
<221> CDS
<222> (140)..(889)
<223> G1307

<400> 39
cccttattgg gcntnancgn ccncccgga ggtctagannn tnancgcccg cgtccttctn 60
ccattttacn cncttgngc ccacccttgt atntcntttt ntngtgntn tttttcntga 120
gggggcaacg gaaaaaaga atg gga aga gca cca tgt tgt gag aaa atg ggg 172
Met Gly Arg Ala Pro Cys Cys Glu Lys Met Gly
1 5 10
gtg aag aga gga cca tgg act cct gaa gaa gat caa atc ttg atc aat 220
Val Lys Arg Gly Pro Trp Thr Pro Glu Glu Asp Gln Ile Leu Ile Asn
15 20 25
tat att cat ctt tat ggt cat tct aat tgg cga gct ctc cca aaa cac 268
Tyr Ile His Leu Tyr Gly His Ser Asn Trp Arg Ala Leu Pro Lys His
30 35 40
gca ggt tta ctt aga tgt ggg aaa agt tgc aga ctt ggt tgg atc aat 316
Ala Gly Leu Leu Arg Cys Gly Lys Ser Cys Arg Leu Gly Trp Ile Asn
45 50 55
tat ctt aga cca gac att aaa cgt ggc aat ttc act cct caa gaa gaa 364
Tyr Leu Arg Pro Asp Ile Lys Arg Gly Asn Phe Thr Pro Gln Glu Glu
60 65 70 75
caa act att atc aat ctg cat gaa agc tta ggc aac aga tgg tct gcg 412

mbi19 Sequence Listing.ST25

Gln Thr Ile Ile Asn Leu His Glu Ser Leu Gly Asn Arg Trp Ser Ala
80 85 90

att gct gca aaa ttg ccg gga cga acc gac aat gaa ata aaa aat gtt 460
Ile Ala Ala Lys Leu Pro Gly Arg Thr Asp Asn Glu Ile Lys Asn Val
95 100 105

tgg cac act cat ttg aag aaa aga ctc agc aaa aat cta aac aat ggc 508
Trp His Thr His Leu Lys Lys Arg Leu Ser Lys Asn Leu Asn Asn Gly
110 115 120

gga gac acc aaa gac gtt aac gga att aac gag acc aca aat gaa gac 556
Gly Asp Thr Lys Asp Val Asn Gly Ile Asn Glu Thr Thr Asn Glu Asp
125 130 135

aaa gga tct gtg ata gtc gac aca gcc tct tta caa caa ttt tct aat 604
Lys Gly Ser Val Ile Val Asp Thr Ala Ser Leu Gln Gln Phe Ser Asn
140 145 150 155

agt att aca aca ttt gat att tca aat gat aac aag gac gat att atg 652
Ser Ile Thr Thr Phe Asp Ile Ser Asn Asp Asn Lys Asp Asp Ile Met
160 165 170

tcg tac gag gat att tct gcc ttg ata gat gat agt ttt tgg tcg gac 700
Ser Tyr Glu Asp Ile Ser Ala Leu Ile Asp Asp Ser Phe Trp Ser Asp
175 180 185

gtc ata tcg gta gat aat tcg aat aag aat gag aag aag ata gag gat 748
Val Ile Ser Val Asp Asn Ser Asn Lys Asn Glu Lys Lys Ile Glu Asp
190 195 200

tgg gaa gga ttg atc gat aga aat agt aaa aaa tgt agc tat agt aat 796
Trp Glu Gly Leu Ile Asp Arg Asn Ser Lys Lys Cys Ser Tyr Ser Asn
205 210 215

tct aag ttg tat aat gat gac atg gag ttt tgg ttt gat gtt ttc act 844
Ser Lys Leu Tyr Asn Asp Asp Met Glu Phe Trp Phe Asp Val Phe Thr
220 225 230 235

agt aat cgt aga att gag gaa ttt tcc gac ata ccc gag ttt taa 889
Ser Asn Arg Arg Ile Glu Glu Phe Ser Asp Ile Pro Glu Phe
240 245

ttttgatttt gattttgtgt tgtttttgtc gttaagactt tgaaagtctt tttgtaatcc 949

aaatgaataa attccttttc tttttaaaaa aaaaaaaaaa aaaaa 994

<210> 40
<211> 249
<212> PRT
<213> Arabidopsis thaliana

<400> 40

Met Gly Arg Ala Pro Cys Cys Glu Lys Met Gly Val Lys Arg Gly Pro
1 5 10 15

Trp Thr Pro Glu Glu Asp Gln Ile Leu Ile Asn Tyr Ile His Leu Tyr
20 25 30

Gly His Ser Asn Trp Arg Ala Leu Pro Lys His Ala Gly Leu Leu Arg
35 40 45

Cys Gly Lys Ser Cys Arg Leu Gly Trp Ile Asn Tyr Leu Arg Pro Asp
50 55 60

Ile Lys Arg Gly Asn Phe Thr Pro Gln Glu Glu Gln Thr Ile Ile Asn
65 70 75 80

mbi19 Sequence Listing.ST25

Leu His Glu Ser Leu Gly Asn Arg Trp Ser Ala Ile Ala Ala Lys Leu
 85 90 95

Pro Gly Arg Thr Asp Asn Glu Ile Lys Asn Val Trp His Thr His Leu
 100 105 110

Lys Lys Arg Leu Ser Lys Asn Leu Asn Asn Gly Gly Asp Thr Lys Asp
 115 120 125

Val Asn Gly Ile Asn Glu Thr Thr Asn Glu Asp Lys Gly Ser Val Ile
 130 135 140

Val Asp Thr Ala Ser Leu Gln Gln Phe Ser Asn Ser Ile Thr Thr Phe
 145 150 155 160

Asp Ile Ser Asn Asp Asn Lys Asp Asp Ile Met Ser Tyr Glu Asp Ile
 165 170 175

Ser Ala Leu Ile Asp Asp Ser Phe Trp Ser Asp Val Ile Ser Val Asp
 180 185 190

Asn Ser Asn Lys Asn Glu Lys Lys Ile Glu Asp Trp Glu Gly Leu Ile
 195 200 205

Asp Arg Asn Ser Lys Lys Cys Ser Tyr Ser Asn Ser Lys Leu Tyr Asn
 210 215 220

Asp Asp Met Glu Phe Trp Phe Asp Val Phe Thr Ser Asn Arg Arg Ile
 225 230 235 240

Glu Glu Phe Ser Asp Ile Pro Glu Phe
 245

<210> 41
 <211> 891
 <212> DNA
 <213> Arabidopsis thaliana

<220>
 <221> CDS
 <222> (1)..(891)
 <223> G1327

<400> 41
 atg ggg aaa gga aga gca cca tgc tgc gac aag aac aaa gtg aag aga 48
 Met Gly Lys Gly Arg Ala Pro Cys Cys Asp Lys Asn Lys Val Lys Arg
 1 5 10 15

ggg cca tgg agc cct caa gaa gat ctc act ctc atc act ttt att caa 96
 Gly Pro Trp Ser Pro Gln Glu Asp Leu Thr Leu Ile Thr Phe Ile Gln
 20 25 30

aaa cat ggc cat caa aac tgg aga tct ctt ccc aag ctt gct gga ttg 144
 Lys His Gly His Gln Asn Trp Arg Ser Leu Pro Lys Leu Ala Gly Leu
 35 40 45

ttg aga tgt ggg aaa agt tgc cga cta aga tgg ata aac tat ctg aga 192
 Leu Arg Cys Gly Lys Ser Cys Arg Leu Arg Trp Ile Asn Tyr Leu Arg
 50 55 60

ccg gac gtg aag cga ggc aac ttt agc aaa aag gag gaa gat gct atc 240
 Pro Asp Val Lys Arg Gly Asn Phe Ser Lys Lys Glu Glu Asp Ala Ile
 65 70 75 80

mbi19 Sequence Listing.ST25

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att cac tac cat caa acc ctt gga aac aag tgg tca aag atc gcg tcc      288
Ile His Tyr His Gln Thr Leu Gly Asn Lys Trp Ser Lys Ile Ala Ser
      85                      90                      95

ttc ttg ccg gga aga act gac aac gag atc aaa aac gtg tgg aac acg      336
Phe Leu Pro Gly Arg Thr Asp Asn Glu Ile Lys Asn Val Trp Asn Thr
      100                    105                    110

cat ctc aag aaa cga ctc act cca tct tct tct tct tca tcc ctc tct      384
His Leu Lys Lys Arg Leu Thr Pro Ser Ser Ser Ser Ser Ser Leu Ser
      115                    120                    125

agc act cat gac caa agc aca aaa gca gat cat gac aag aac tgt gac      432
Ser Thr His Asp Gln Ser Thr Lys Ala Asp His Asp Lys Asn Cys Asp
      130                    135                    140

ggg gct caa gaa gaa ata cat tca ggg tta aat gag agc caa aac tca      480
Gly Ala Gln Glu Glu Ile His Ser Gly Leu Asn Glu Ser Gln Asn Ser
      145                    150                    155                    160

gct act tcg tca cat cac caa ggc gag tgt atg cac aca aaa cca gag      528
Ala Thr Ser Ser His His Gln Gly Glu Cys Met His Thr Lys Pro Glu
      165                    170                    175

ctt cat gag gtt aat gga ctc aac gag atc cag ttc ctg ctc gac cat      576
Leu His Glu Val Asn Gly Leu Asn Glu Ile Gln Phe Leu Leu Asp His
      180                    185                    190

gat gac ttt gat gat ata acc tct gag ttt ctt cag gat aac gat atc      624
Asp Asp Phe Asp Asp Ile Thr Ser Glu Phe Leu Gln Asp Asn Asp Ile
      195                    200                    205

tta ttt ccg cta gac tct ctt ctt cat aac cac caa act cac att tca      672
Leu Phe Pro Leu Asp Ser Leu Leu His Asn His Gln Thr His Ile Ser
      210                    215                    220

acc caa gaa atg act cga gag gta acc aaa tcg caa tca ttt gat cat      720
Thr Gln Glu Met Thr Arg Glu Val Thr Lys Ser Gln Ser Phe Asp His
      225                    230                    235                    240

cct caa ccg gat atc cca tgc gga ttt gaa gac aca aac gaa gaa tcc      768
Pro Gln Pro Asp Ile Pro Cys Gly Phe Glu Asp Thr Asn Glu Glu Ser
      245                    250                    255

gac ttg agg aga cag ctg gtt gaa tca acc aca cct aac aat gag tac      816
Asp Leu Arg Arg Gln Leu Val Glu Ser Thr Thr Pro Asn Asn Glu Tyr
      260                    265                    270

gac gag tgg ttc aac ttc att gac aac caa act tac ttt gat gat ttt      864
Asp Glu Trp Phe Asn Phe Ile Asp Asn Gln Thr Tyr Phe Asp Asp Phe
      275                    280                    285

aat ttc gtc gga gaa gta tgt cta tga      891
Asn Phe Val Gly Glu Val Cys Leu
      290                    295

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<210> 42
<211> 296
<212> PRT
<213> Arabidopsis thaliana

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<400> 42

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Met Gly Lys Gly Arg Ala Pro Cys Cys Asp Lys Asn Lys Val Lys Arg
1                      5                      10                      15

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Gly Pro Trp Ser Pro Gln Glu Asp Leu Thr Leu Ile Thr Phe Ile Gln
20                      25                      30

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Lys His Gly His Gln Asn Trp Arg Ser Leu Pro Lys Leu Ala Gly Leu
35                      40                      45

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mbi19 Sequence Listing.ST25

Leu Arg Cys Gly Lys Ser Cys Arg Leu Arg Trp Ile Asn Tyr Leu Arg
50 55 60

Pro Asp Val Lys Arg Gly Asn Phe Ser Lys Lys Glu Glu Asp Ala Ile
65 70 75 80

Ile His Tyr His Gln Thr Leu Gly Asn Lys Trp Ser Lys Ile Ala Ser
85 90 95

Phe Leu Pro Gly Arg Thr Asp Asn Glu Ile Lys Asn Val Trp Asn Thr
100 105 110

His Leu Lys Lys Arg Leu Thr Pro Ser Ser Ser Ser Ser Ser Leu Ser
115 120 125

Ser Thr His Asp Gln Ser Thr Lys Ala Asp His Asp Lys Asn Cys Asp
130 135 140

Gly Ala Gln Glu Glu Ile His Ser Gly Leu Asn Glu Ser Gln Asn Ser
145 150 155 160

Ala Thr Ser Ser His His Gln Gly Glu Cys Met His Thr Lys Pro Glu
165 170 175

Leu His Glu Val Asn Gly Leu Asn Glu Ile Gln Phe Leu Leu Asp His
180 185 190

Asp Asp Phe Asp Asp Ile Thr Ser Glu Phe Leu Gln Asp Asn Asp Ile
195 200 205

Leu Phe Pro Leu Asp Ser Leu Leu His Asn His Gln Thr His Ile Ser
210 215 220

Thr Gln Glu Met Thr Arg Glu Val Thr Lys Ser Gln Ser Phe Asp His
225 230 235 240

Pro Gln Pro Asp Ile Pro Cys Gly Phe Glu Asp Thr Asn Glu Glu Ser
245 250 255

Asp Leu Arg Arg Gln Leu Val Glu Ser Thr Thr Pro Asn Asn Glu Tyr
260 265 270

Asp Glu Trp Phe Asn Phe Ile Asp Asn Gln Thr Tyr Phe Asp Asp Phe
275 280 285

Asn Phe Val Gly Glu Val Cys Leu
290 295

<210> 43
<211> 1237
<212> DNA
<213> Arabidopsis thaliana

<220>
<221> CDS
<222> (73)..(954)
<223> G673

mbi19 Sequence Listing.ST25

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<400> 43
tctctctcta accccttctc tcttcagtct ctctctctct agacgatctc tatcttgaat      60

aaaataccga ta atg acc tca acc aat ccg gtg gtc gcc gaa gta ata ccg      111
      Met Thr Ser Thr Asn Pro Val Val Ala Glu Val Ile Pro
      1                    5                    10

gcg gaa act tct aca gat gct aca gag acg acg att gca acg acg gaa      159
Ala Glu Thr Ser Thr Asp Ala Thr Glu Thr Thr Ile Ala Thr Thr Glu
      15                    20                    25

gct ggt gaa gca ccg gag aag aag gtg agg aaa gct tat aca atc acc      207
Ala Gly Glu Ala Pro Glu Lys Lys Val Arg Lys Ala Tyr Thr Ile Thr
      30                    35                    40                    45

aag tct aga gag agt tgg act gaa gga gaa cac gac aag ttt ctg gaa      255
Lys Ser Arg Glu Ser Trp Thr Glu Gly Glu His Asp Lys Phe Leu Glu
      50                    55                    60

gct ctt caa ttg ttt gat cgt gac tgg aaa aag ata gaa gat ttt gtt      303
Ala Leu Gln Leu Phe Asp Arg Asp Trp Lys Lys Ile Glu Asp Phe Val
      65                    70                    75

ggg tca aag aca gtt att cag atc agg agc cat gcc caa aaa tac ttt      351
Gly Ser Lys Thr Val Ile Gln Ile Arg Ser His Ala Gln Lys Tyr Phe
      80                    85                    90

cta aag gtc caa aaa aat ggg act tta gca cat gtt cca ccc cct agg      399
Leu Lys Val Gln Lys Asn Gly Thr Leu Ala His Val Pro Pro Pro Arg
      95                    100                    105

cct aag cgc aaa gct gct cat cca tat cct caa aag gca tcg aaa aat      447
Pro Lys Arg Lys Ala Ala His Pro Tyr Pro Gln Lys Ala Ser Lys Asn
      110                    115                    120                    125

gct caa atg tcg ctt cac gtt tcc atg tcc ttt cct act caa ata aat      495
Ala Gln Met Ser Leu His Val Ser Met Ser Phe Pro Thr Gln Ile Asn
      130                    135                    140

aac ctg cct gga tat act cca tgg gat gat gat aca tct gca ttg tta      543
Asn Leu Pro Gly Tyr Thr Pro Trp Asp Asp Asp Thr Ser Ala Leu Leu
      145                    150                    155

aac att gct gta agt ggg gtt att cca cca gaa gat gaa ctt gat act      591
Asn Ile Ala Val Ser Gly Val Ile Pro Pro Glu Asp Glu Leu Asp Thr
      160                    165                    170

ctt tgt gga gca gaa gtt gat gtt gga tca aat gac atg ata agt gaa      639
Leu Cys Gly Ala Glu Val Asp Val Gly Ser Asn Asp Met Ile Ser Glu
      175                    180                    185

act agt cct tca gca tct ggt atc gga agc tca agc aga aca cta tca      687
Thr Ser Pro Ser Ala Ser Gly Ile Gly Ser Ser Ser Arg Thr Leu Ser
      190                    195                    200                    205

gat tct aag ggt ttg aga ctg gcg aaa caa gct ccc tca atg cat ggt      735
Asp Ser Lys Gly Leu Arg Leu Ala Lys Gln Ala Pro Ser Met His Gly
      210                    215                    220

ctt cct gat ttt gct gag gtt tat aac ttc att ggg agt gtg ttc gat      783
Leu Pro Asp Phe Ala Glu Val Tyr Asn Phe Ile Gly Ser Val Phe Asp
      225                    230                    235

cct gac agc aaa ggc cgc atg aaa aag ctc aag gaa atg gat cct ata      831
Pro Asp Ser Lys Gly Arg Met Lys Lys Leu Lys Glu Met Asp Pro Ile
      240                    245                    250

aat ttc gaa act gtt ttg ctg ttg atg aga aac ctc aca gtg aac ttg      879
Asn Phe Glu Thr Val Leu Leu Leu Met Arg Asn Leu Thr Val Asn Leu
      255                    260                    265

tca aac cct gac ttt gaa cct act tct gaa tat gtt gat gct gca gag      927
Ser Asn Pro Asp Phe Glu Pro Thr Ser Glu Tyr Val Asp Ala Ala Glu

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270	275	280	285	
gaa ggt cat gaa cac tta agc tct tag ctgtttgtgc actcaacaag				974
Glu Gly His Glu His Leu Ser Ser				
290				
ttatatatct tcttgacgac ttcttgctcg caacaactct ctaccagcta tcaaattgcat				1034
cgtacgggttg ttgtctgagg agaacataac tgagtcgctcg tcacaaacaa gaggaacata				1094
tgcagtttctg gtcagaacca gtcgtgtgaa tggtagatat atgtatgtgt gtgtagaaaa				1154
tggttaccaa ttgtatcttc tttttgataa ttatttttttc atgccttttg taatatgtaa				1214
gtttcttttaa aaaaaaaaaa aaa				1237
<210> 44				
<211> 293				
<212> PRT				
<213> Arabidopsis thaliana				
<400> 44				
Met Thr Ser Thr Asn Pro Val Val Ala Glu Val Ile Pro Ala Glu Thr				
1 5 10 15				
Ser Thr Asp Ala Thr Glu Thr Thr Ile Ala Thr Thr Glu Ala Gly Glu				
20 25 30				
Ala Pro Glu Lys Lys Val Arg Lys Ala Tyr Thr Ile Thr Lys Ser Arg				
35 40 45				
Glu Ser Trp Thr Glu Gly Glu His Asp Lys Phe Leu Glu Ala Leu Gln				
50 55 60				
Leu Phe Asp Arg Asp Trp Lys Lys Ile Glu Asp Phe Val Gly Ser Lys				
65 70 75 80				
Thr Val Ile Gln Ile Arg Ser His Ala Gln Lys Tyr Phe Leu Lys Val				
85 90 95				
Gln Lys Asn Gly Thr Leu Ala His Val Pro Pro Pro Arg Pro Lys Arg				
100 105 110				
Lys Ala Ala His Pro Tyr Pro Gln Lys Ala Ser Lys Asn Ala Gln Met				
115 120 125				
Ser Leu His Val Ser Met Ser Phe Pro Thr Gln Ile Asn Asn Leu Pro				
130 135 140				
Gly Tyr Thr Pro Trp Asp Asp Asp Thr Ser Ala Leu Leu Asn Ile Ala				
145 150 155 160				
Val Ser Gly Val Ile Pro Pro Glu Asp Glu Leu Asp Thr Leu Cys Gly				
165 170 175				
Ala Glu Val Asp Val Gly Ser Asn Asp Met Ile Ser Glu Thr Ser Pro				
180 185 190				
Ser Ala Ser Gly Ile Gly Ser Ser Ser Arg Thr Leu Ser Asp Ser Lys				
195 200 205				

mbil9 Sequence Listing.ST25

Gly Leu Arg Leu Ala Lys Gln Ala Pro Ser Met His Gly Leu Pro Asp
 210 215 220

Phe Ala Glu Val Tyr Asn Phe Ile Gly Ser Val Phe Asp Pro Asp Ser
 225 230 235 240

Lys Gly Arg Met Lys Lys Leu Lys Glu Met Asp Pro Ile Asn Phe Glu
 245 250 255

Thr Val Leu Leu Leu Met Arg Asn Leu Thr Val Asn Leu Ser Asn Pro
 260 265 270

Asp Phe Glu Pro Thr Ser Glu Tyr Val Asp Ala Ala Glu Glu Gly His
 275 280 285

Glu His Leu Ser Ser
 290

<210> 45
 <211> 1764
 <212> DNA
 <213> Arabidopsis thaliana

<220>
 <221> CDS
 <222> (1)..(1764)
 <223> G307

<400> 45
 atg aag aga gat cat cac caa ttc caa ggt cga ttg tcc aac cac ggg 48
 Met Lys Arg Asp His His Gln Phe Gln Gly Arg Leu Ser Asn His Gly
 1 5 10 15

act tct tct tct tca tca tca atc tct aaa gat aag atg atg atg gtg 96
 Thr Ser Ser Ser Ser Ser Ser Ile Ser Lys Asp Lys Met Met Met Val
 20 25 30

aaa aaa gaa gaa gac ggt gga ggt aac atg gac gac gag ctt ctc gct 144
 Lys Lys Glu Glu Asp Gly Gly Gly Asn Met Asp Asp Glu Leu Leu Ala
 35 40 45

gtt tta ggt tac aaa gtt agg tca tcg gag atg gcg gag gtt gct ttg 192
 Val Leu Gly Tyr Lys Val Arg Ser Ser Glu Met Ala Glu Val Ala Leu
 50 55 60

aaa ctc gaa caa tta gag acg atg atg agt aat gtt caa gaa gat ggt 240
 Lys Leu Glu Gln Leu Thr Thr Met Met Ser Asn Val Gln Glu Asp Gly
 65 70 75 80

tta tct cat ctc gcg acg gat act gtt cat tat aat ccg tcg gag ctt 288
 Leu Ser His Leu Ala Thr Asp Thr Val His Tyr Asn Pro Ser Glu Leu
 85 90 95

tat tct tgg ctt gat aat atg ctc tct gag ctt aat cct cct cct ctt 336
 Tyr Ser Trp Leu Asp Asn Met Leu Ser Glu Leu Asn Pro Pro Pro Leu
 100 105 110

ccg gcg agt tct aac ggt tta gat ccg gtt ctt cct tcg ccg gag att 384
 Pro Ala Ser Ser Asn Gly Leu Asp Pro Val Leu Pro Ser Pro Glu Ile
 115 120 125

tgt ggt ttt ccg gct tcg gat tat gac ctt aaa gtc att ccc gga aac 432
 Cys Gly Phe Pro Ala Ser Asp Tyr Asp Leu Lys Val Ile Pro Gly Asn
 130 135 140

gcg att tat cag ttt ccg gcg att gat tct tcg tct tcg tcg aat aat 480
 Ala Ile Tyr Gln Phe Pro Ala Ile Asp Ser Ser Ser Ser Ser Asn Asn

mbi19 Sequence Listing.ST25

145	150	155	160	
cag aac aag cgt ttg	aaa tca tgc tcg agt cct gat tct atg gtt aca			528
Gln Asn Lys Arg Leu	Lys Ser Cys Ser Ser Pro Asp Ser Met Val Thr			
	165	170	175	
tcg act tcg acg ggt acg cag att ggt gga gtc ata gga acg acg gtg				576
Ser Thr Ser Thr Gly Thr Gln Ile Gly Gly Val Ile Gly Thr Thr Val				
	180	185	190	
acg aca acc acc acg aca acg acg gcg gcg gct gag tca act cgt tct				624
Thr Thr Thr Thr Thr Thr Thr Thr Ala Ala Ala Glu Ser Thr Arg Ser				
	195	200	205	
gtt atc ctg gtt gac tcg caa gag aac ggt gtt cgt tta gtc cac gcg				672
Val Ile Leu Val Asp Ser Gln Glu Asn Gly Val Arg Leu Val His Ala				
	210	215	220	
ctt atg gct tgt gca gaa gca atc cag cag aac aat ttg act cta gcg				720
Leu Met Ala Cys Ala Glu Ala Ile Gln Gln Asn Asn Leu Thr Leu Ala				
	225	230	235	240
gaa gct ctt gtg aag caa atc gga tgc tta gct gtg tct caa gcc gga				768
Glu Ala Leu Val Lys Gln Ile Gly Cys Leu Ala Val Ser Gln Ala Gly				
	245	250	255	
gct atg aga aaa gtg gct act tac ttc gcc gaa gct tta gct cgg cgg				816
Ala Met Arg Lys Val Ala Thr Tyr Phe Ala Glu Ala Leu Ala Arg Arg				
	260	265	270	
atc tac cgt ctc tct ccg ccg cag aat cag atc gat cat tgt ctc tcc				864
Ile Tyr Arg Leu Ser Pro Pro Gln Asn Gln Ile Asp His Cys Leu Ser				
	275	280	285	
gat act ctt cag atg cac ttt tac gag act tgt cct tat ctt aaa ttc				912
Asp Thr Leu Gln Met His Phe Tyr Glu Thr Cys Pro Tyr Leu Lys Phe				
	290	295	300	
gct cac ttc acg gcg aac caa gcg att ctc gaa gct ttt gaa ggt aag				960
Ala His Phe Thr Ala Asn Gln Ala Ile Leu Glu Ala Phe Glu Gly Lys				
	305	310	315	320
aag aga gta cac gtc att gat ttc tcg atg aac caa ggt ctt caa tgg				1008
Lys Arg Val His Val Ile Asp Phe Ser Met Asn Gln Gly Leu Gln Trp				
	325	330	335	
cct gcg ctt atg caa gct ctt gcg ctt cga gaa gga ggt cct cca act				1056
Pro Ala Leu Met Gln Ala Leu Ala Leu Arg Glu Gly Gly Pro Pro Thr				
	340	345	350	
ttc cgg tta acc gga att ggt cca ccg gcg ccg gat aat tct gat cat				1104
Phe Arg Leu Thr Gly Ile Gly Pro Pro Ala Pro Asp Asn Ser Asp His				
	355	360	365	
ctt cat gaa gtt ggt tgt aaa tta gct cag ctt gcg gag gcg att cac				1152
Leu His Glu Val Gly Cys Lys Leu Ala Gln Leu Ala Glu Ala Ile His				
	370	375	380	
gta gaa ttc gaa tac cgt gga ttc gtt gct aac agc tta gcc gat ctc				1200
Val Glu Phe Glu Tyr Arg Gly Phe Val Ala Asn Ser Leu Ala Asp Leu				
	385	390	395	400
gat gct tcg atg ctt gag ctt aga ccg agc gat acg gaa gct gtt gcg				1248
Asp Ala Ser Met Leu Glu Leu Arg Pro Ser Asp Thr Glu Ala Val Ala				
	405	410	415	
gtg aac tct gtt ttt gag cta cat aag ctc tta ggt cgt ccc ggt ggg				1296
Val Asn Ser Val Phe Glu Leu His Lys Leu Leu Gly Arg Pro Gly Gly				
	420	425	430	
ata gag aaa gtt ctc ggc gtt gtg aaa cag att aaa ccg gtg att ttc				1344
Ile Glu Lys Val Leu Gly Val Val Lys Gln Ile Lys Pro Val Ile Phe				
	435	440	445	
acg gtg gtt gag caa gaa tcg aac cat aac gga ccg gtt ttc tta gac				1392

mbi19 Sequence Listing.ST25

Thr Val Val Glu Gln Glu Ser Asn His Asn Gly Pro Val Phe Leu Asp
 450 455 460

cgg ttt act gaa tcg tta cat tat tat tcg act ctg ttt gat tcg ttg 1440
 Arg Phe Thr Glu Ser Leu His Tyr Tyr Ser Thr Leu Phe Asp Ser Leu
 465 470 475 480

gaa gga gtt ccg aat agt caa gac aaa gtc atg tct gaa gtt tac tta 1488
 Glu Gly Val Pro Asn Ser Gln Asp Lys Val Met Ser Glu Val Tyr Leu
 485 490 495

ggg aaa cag att tgt aat ctg gtg gct tgt gaa ggt cct gac aga gtc 1536
 Gly Lys Gln Ile Cys Asn Leu Val Ala Cys Glu Gly Pro Asp Arg Val
 500 505 510

gag aga cac gaa acg ttg agt caa tgg gga aac cgg ttt ggt tcg tcc 1584
 Glu Arg His Glu Thr Leu Ser Gln Trp Gly Asn Arg Phe Gly Ser Ser
 515 520 525

ggg tta gcg ccg gca cat ctt ggg tct aac gcg ttt aag caa gcg agt 1632
 Gly Leu Ala Pro Ala His Leu Gly Ser Asn Ala Phe Lys Gln Ala Ser
 530 535 540

atg ctt ttg tct gtg ttt aat agt ggc caa ggt tat cgt gtg gag gag 1680
 Met Leu Leu Ser Val Phe Asn Ser Gly Gln Gly Tyr Arg Val Glu Glu
 545 550 555 560

agt aat gga tgt ttg atg ttg ggt tgg cac act cgc cca ctc att acc 1728
 Ser Asn Gly Cys Leu Met Leu Gly Trp His Thr Arg Pro Leu Ile Thr
 565 570 575

acc tcc gct tgg aaa ctc tcg acg gcg gcg cac tga 1764
 Thr Ser Ala Trp Lys Leu Ser Thr Ala Ala His
 580 585

<210> 46
 <211> 587
 <212> PRT
 <213> Arabidopsis thaliana

<400> 46

Met Lys Arg Asp His His Gln Phe Gln Gly Arg Leu Ser Asn His Gly
 1 5 10 15

Thr Ser Ser Ser Ser Ser Ser Ile Ser Lys Asp Lys Met Met Met Val
 20 25 30

Lys Lys Glu Glu Asp Gly Gly Gly Asn Met Asp Asp Glu Leu Leu Ala
 35 40 45

Val Leu Gly Tyr Lys Val Arg Ser Ser Glu Met Ala Glu Val Ala Leu
 50 55 60

Lys Leu Glu Gln Leu Glu Thr Met Met Ser Asn Val Gln Glu Asp Gly
 65 70 75 80

Leu Ser His Leu Ala Thr Asp Thr Val His Tyr Asn Pro Ser Glu Leu
 85 90 95

Tyr Ser Trp Leu Asp Asn Met Leu Ser Glu Leu Asn Pro Pro Pro Leu
 100 105 110

Pro Ala Ser Ser Asn Gly Leu Asp Pro Val Leu Pro Ser Pro Glu Ile
 115 120 125

mbil9 Sequence Listing.ST25

Cys Gly Phe Pro Ala Ser Asp Tyr Asp Leu Lys Val Ile Pro Gly Asn
 130 135 140

Ala Ile Tyr Gln Phe Pro Ala Ile Asp Ser Ser Ser Ser Ser Asn Asn
 145 150 155 160

Gln Asn Lys Arg Leu Lys Ser Cys Ser Ser Pro Asp Ser Met Val Thr
 165 170 175

Ser Thr Ser Thr Gly Thr Gln Ile Gly Gly Val Ile Gly Thr Thr Val
 180 185 190

Thr Thr Thr Thr Thr Thr Thr Thr Ala Ala Ala Glu Ser Thr Arg Ser
 195 200 205

Val Ile Leu Val Asp Ser Gln Glu Asn Gly Val Arg Leu Val His Ala
 210 215 220

Leu Met Ala Cys Ala Glu Ala Ile Gln Gln Asn Asn Leu Thr Leu Ala
 225 230 235 240

Glu Ala Leu Val Lys Gln Ile Gly Cys Leu Ala Val Ser Gln Ala Gly
 245 250 255

Ala Met Arg Lys Val Ala Thr Tyr Phe Ala Glu Ala Leu Ala Arg Arg
 260 265 270

Ile Tyr Arg Leu Ser Pro Pro Gln Asn Gln Ile Asp His Cys Leu Ser
 275 280 285

Asp Thr Leu Gln Met His Phe Tyr Glu Thr Cys Pro Tyr Leu Lys Phe
 290 295 300

Ala His Phe Thr Ala Asn Gln Ala Ile Leu Glu Ala Phe Glu Gly Lys
 305 310 315 320

Lys Arg Val His Val Ile Asp Phe Ser Met Asn Gln Gly Leu Gln Trp
 325 330 335

Pro Ala Leu Met Gln Ala Leu Ala Leu Arg Glu Gly Gly Pro Pro Thr
 340 345 350

Phe Arg Leu Thr Gly Ile Gly Pro Pro Ala Pro Asp Asn Ser Asp His
 355 360 365

Leu His Glu Val Gly Cys Lys Leu Ala Gln Leu Ala Glu Ala Ile His
 370 375 380

Val Glu Phe Glu Tyr Arg Gly Phe Val Ala Asn Ser Leu Ala Asp Leu
 385 390 395 400

Asp Ala Ser Met Leu Glu Leu Arg Pro Ser Asp Thr Glu Ala Val Ala
 405 410 415

Val Asn Ser Val Phe Glu Leu His Lys Leu Leu Gly Arg Pro Gly Gly
 420 425 430

mbil9 Sequence Listing.ST25

Ile Glu Lys Val Leu Gly Val Val Lys Gln Ile Lys Pro Val Ile Phe
435 440 445

Thr Val Val Glu Gln Glu Ser Asn His Asn Gly Pro Val Phe Leu Asp
450 455 460

Arg Phe Thr Glu Ser Leu His Tyr Tyr Ser Thr Leu Phe Asp Ser Leu
465 470 475 480

Glu Gly Val Pro Asn Ser Gln Asp Lys Val Met Ser Glu Val Tyr Leu
485 490 495

Gly Lys Gln Ile Cys Asn Leu Val Ala Cys Glu Gly Pro Asp Arg Val
500 505 510

Glu Arg His Glu Thr Leu Ser Gln Trp Gly Asn Arg Phe Gly Ser Ser
515 520 525

Gly Leu Ala Pro Ala His Leu Gly Ser Asn Ala Phe Lys Gln Ala Ser
530 535 540

Met Leu Leu Ser Val Phe Asn Ser Gly Gln Gly Tyr Arg Val Glu Glu
545 550 555 560

Ser Asn Gly Cys Leu Met Leu Gly Trp His Thr Arg Pro Leu Ile Thr
565 570 575

Thr Ser Ala Trp Lys Leu Ser Thr Ala Ala His
580 585

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<212> DNA
<213> Arabidopsis thaliana

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<222> (7)..(813)
<223> G529

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aag tta gcc gag caa gct gag cgt tac gag gaa atg gtt gag ttc atg 96
Lys Leu Ala Glu Gln Ala Glu Arg Tyr Glu Glu Met Val Glu Phe Met
15 20 25 30

gag aaa gtt gca aag acc gtg gag acc gag gaa ctt act gtt gaa gag 144
Glu Lys Val Ala Lys Thr Val Glu Thr Glu Glu Leu Thr Val Glu Glu
35 40 45

agg aat ctc ttg tct gtt gct tac aag aac gtg att ggt gct agg aga 192
Arg Asn Leu Leu Ser Val Ala Tyr Lys Asn Val Ile Gly Ala Arg Arg
50 55 60

gct tct tgg agg att atc tct tcc att gag cag aag gaa gat agc agg 240
Ala Ser Trp Arg Ile Ile Ser Ser Ile Glu Gln Lys Glu Asp Ser Arg
65 70 75

ggc aac agt gat cat gtt tcg att atc aag gat tac aga ggc aag att 288
Gly Asn Ser Asp His Val Ser Ile Ile Lys Asp Tyr Arg Gly Lys Ile

mbi19 Sequence Listing.ST25

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gct cat ctc att cct gct gct tct ttg gct gag tcc aaa gtt ttt tac Ala His Leu Ile Pro Ala Ala Ser Leu Ala Glu Ser Lys Val Phe Tyr 115 120 125			384
ctg aag atg aag gga gat tat cat cgg tac ctt gct gaa ttc aag act Leu Lys Met Lys Gly Asp Tyr His Arg Tyr Leu Ala Glu Phe Lys Thr 130 135 140			432
ggg gct gag agg aaa gaa gct gct gag agc act ctt gtt gcc tac aag Gly Ala Glu Arg Lys Glu Ala Ala Glu Ser Thr Leu Val Ala Tyr Lys 145 150 155			480
tct gct cag gat att gct ctt gct gat ctg gct ccc act cac cca atc Ser Ala Gln Asp Ile Ala Leu Ala Asp Leu Ala Pro Thr His Pro Ile 160 165 170			528
aga ctg ggg ctt gct ctt aac ttc tct gtt ttc tac tat gag att ctc Arg Leu Gly Leu Ala Leu Asn Phe Ser Val Phe Tyr Tyr Glu Ile Leu 175 180 185 190			576
aac tca tct gat cgt gcg tgt agt ctc gca aag cag gct ttt gat gag Asn Ser Ser Asp Arg Ala Cys Ser Leu Ala Lys Gln Ala Phe Asp Glu 195 200 205			624
gca atc tcg gag cta gac aca ttg gga gag gaa tca tac aag gac agt Ala Ile Ser Glu Leu Asp Thr Leu Gly Glu Glu Ser Tyr Lys Asp Ser 210 215 220			672
aca ttg atc atg cag ctt ctc cgt gac aat ctc acc ctc tgg act tct Thr Leu Ile Met Gln Leu Leu Arg Asp Asn Leu Thr Leu Trp Thr Ser 225 230 235			720
gac ctc aat gac gaa gct ggt gat gat atc aag gaa gcc ccg aaa gag Asp Leu Asn Asp Glu Ala Gly Asp Asp Ile Lys Glu Ala Pro Lys Glu 240 245 250			768
gtg cag aaa gtt gat gaa caa gcc caa cca cca cct tcg cag tga Val Gln Lys Val Asp Glu Gln Ala Gln Pro Pro Pro Ser Gln 255 260 265			813
taaaatcaga tccatggaat gatttgcaga caaaaagata tatggcttgg ttctgtgttt			873
ttaaacagaa aaaaaccttg tagtttcctt aaacatgggc tgtagtttcc ttaaacatgg			933
atgtgtagta gtaattgtag ctgcatgatt tggttatoga tggttaaaaa aaaaaaa			990
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Ala Glu Gln Ala Glu Arg Tyr Glu Glu Met Val Glu Phe Met Glu Lys 20 25 30			
Val Ala Lys Thr Val Glu Thr Glu Glu Leu Thr Val Glu Glu Arg Asn 35 40 45			
Leu Leu Ser Val Ala Tyr Lys Asn Val Ile Gly Ala Arg Arg Ala Ser 50 55 60			

mbil9 Sequence Listing.ST25

Trp Arg Ile Ile Ser Ser Ile Glu Gln Lys Glu Asp Ser Arg Gly Asn
65 70 75 80

Ser Asp His Val Ser Ile Ile Lys Asp Tyr Arg Gly Lys Ile Glu Thr
85 90 95

Glu Leu Ser Lys Ile Cys Asp Gly Ile Leu Asn Leu Leu Glu Ala His
100 105 110

Leu Ile Pro Ala Ala Ser Leu Ala Glu Ser Lys Val Phe Tyr Leu Lys
115 120 125

Met Lys Gly Asp Tyr His Arg Tyr Leu Ala Glu Phe Lys Thr Gly Ala
130 135 140

Glu Arg Lys Glu Ala Ala Glu Ser Thr Leu Val Ala Tyr Lys Ser Ala
145 150 155 160

Gln Asp Ile Ala Leu Ala Asp Leu Ala Pro Thr His Pro Ile Arg Leu
165 170 175

Gly Leu Ala Leu Asn Phe Ser Val Phe Tyr Tyr Glu Ile Leu Asn Ser
180 185 190

Ser Asp Arg Ala Cys Ser Leu Ala Lys Gln Ala Phe Asp Glu Ala Ile
195 200 205

Ser Glu Leu Asp Thr Leu Gly Glu Glu Ser Tyr Lys Asp Ser Thr Leu
210 215 220

Ile Met Gln Leu Leu Arg Asp Asn Leu Thr Leu Trp Thr Ser Asp Leu
225 230 235 240

Asn Asp Glu Ala Gly Asp Asp Ile Lys Glu Ala Pro Lys Glu Val Gln
245 250 255

Lys Val Asp Glu Gln Ala Gln Pro Pro Pro Ser Gln
260 265

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<223> G531

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Met Ser Ser Ser Arg Glu Glu Asn Val Tyr Leu Ala Lys Leu
1 5 10

gct gag caa gct gaa cgt tat gag gaa atg gtt gag ttc atg gag aaa 159
Ala Glu Gln Ala Glu Arg Tyr Glu Glu Met Val Glu Phe Met Glu Lys
15 20 25 30

mbil9 Sequence Listing.ST25

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tgg agg atc ata tct tcc att gaa cag aag gaa gaa agc aga gga aac Trp Arg Ile Ile Ser Ser Ile Glu Gln Lys Glu Glu Ser Arg Gly Asn 65 70 75	303
gat gat cat gtt tcc att atc aag gac tac aga gga aag atc gaa act Asp Asp His Val Ser Ile Ile Lys Asp Tyr Arg Gly Lys Ile Glu Thr 80 85 90	351
gaa ctc agc aaa atc tgt gat gga ata ctc aat ctt ctg gat tct cac Glu Leu Ser Lys Ile Cys Asp Gly Ile Leu Asn Leu Asp Ser His 95 100 105 110	399
ctt gtt ccc act gca tct ttg gcc gag tcc aaa gtc ttt tac ctc aaa Leu Val Pro Thr Ala Ser Leu Ala Glu Ser Lys Val Phe Tyr Leu Lys 115 120 125	447
atg aaa gga gat tac cac agg tac ctt gct gag ttt aag act gga gct Met Lys Gly Asp Tyr His Arg Tyr Leu Ala Glu Phe Lys Thr Gly Ala 130 135 140	495
gag agg aaa gaa gct gct gag agc act ctg gtt gct tac aag tca gct Glu Arg Lys Glu Ala Ala Glu Ser Thr Leu Val Ala Tyr Lys Ser Ala 145 150 155	543
cag gat att gca ctt gct gat tta gct cct act cat ccg att aga ctg Gln Asp Ile Ala Leu Ala Asp Leu Ala Pro Thr His Pro Ile Arg Leu 160 165 170	591
gga ctt gct ctt aac ttc tct gtc ttc tac tac gag att ctc aac tca Gly Leu Ala Leu Asn Phe Ser Val Phe Tyr Tyr Glu Ile Leu Asn Ser 175 180 185 190	639
cct gat cgt gcc tgc agt ctc gca aaa cag gct ttt gat gag gcc att Pro Asp Arg Ala Cys Ser Leu Ala Lys Gln Ala Phe Asp Glu Ala Ile 195 200 205	687
tct gag ctg gat aca tta gga gaa gaa tca tac aaa gac agt acg ttg Ser Glu Leu Asp Thr Leu Gly Glu Glu Ser Tyr Lys Asp Ser Thr Leu 210 215 220	735
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aat gat gag gcg ggc ggt gat gag atc aag gag gcg tca aaa cat gag Asn Asp Glu Ala Gly Gly Asp Glu Ile Lys Glu Ala Ser Lys His Glu 240 245 250	831
ccg gaa gag ggg aaa cca gct gag aca ggg cag tga ccagagagag Pro Glu Glu Gly Lys Pro Ala Glu Thr Gly Gln 255 260 265	877
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mbil9 Sequence Listing.ST25

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Gln Ala Glu Arg Tyr Glu Glu Met Val Glu Phe Met Glu Lys Val Ala
 20 25 30

Lys Thr Val Asp Thr Asp Glu Leu Thr Val Glu Glu Arg Asn Leu Leu
 35 40 45

Ser Val Ala Tyr Lys Asn Val Ile Gly Ala Arg Arg Ala Ser Trp Arg
 50 55 60

Ile Ile Ser Ser Ile Glu Gln Lys Glu Glu Ser Arg Gly Asn Asp Asp
 65 70 75 80

His Val Ser Ile Ile Lys Asp Tyr Arg Gly Lys Ile Glu Thr Glu Leu
 85 90 95

Ser Lys Ile Cys Asp Gly Ile Leu Asn Leu Leu Asp Ser His Leu Val
 100 105 110

Pro Thr Ala Ser Leu Ala Glu Ser Lys Val Phe Tyr Leu Lys Met Lys
 115 120 125

Gly Asp Tyr His Arg Tyr Leu Ala Glu Phe Lys Thr Gly Ala Glu Arg
 130 135 140

Lys Glu Ala Ala Glu Ser Thr Leu Val Ala Tyr Lys Ser Ala Gln Asp
 145 150 155 160

Ile Ala Leu Ala Asp Leu Ala Pro Thr His Pro Ile Arg Leu Gly Leu
 165 170 175

Ala Leu Asn Phe Ser Val Phe Tyr Tyr Glu Ile Leu Asn Ser Pro Asp
 180 185 190

Arg Ala Cys Ser Leu Ala Lys Gln Ala Phe Asp Glu Ala Ile Ser Glu
 195 200 205

Leu Asp Thr Leu Gly Glu Glu Ser Tyr Lys Asp Ser Thr Leu Ile Met
 210 215 220

Gln Leu Leu Arg Asp Asn Leu Thr Leu Trp Asn Ser Asp Ile Asn Asp
 225 230 235 240

Glu Ala Gly Gly Asp Glu Ile Lys Glu Ala Ser Lys His Glu Pro Glu
 245 250 255

Glu Gly Lys Pro Ala Glu Thr Gly Gln
 260 265

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mbil9 Sequence Listing.ST25

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<222> (238)..(2064)

<223> G214

<400> 51

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gaattaaaaa tggaatcttt atcgaatcca agctgatttt gtttctttca ttgaatcatc      180
tctctaaaagt ggaattttgt aaagagaaga tctgaagttg tgtagaggag cttagtg      237
atg gag aca aat tcg tct gga gaa gat ctg gtt att aag act cgg aag      285
Met Glu Thr Asn Ser Ser Gly Glu Asp Leu Val Ile Lys Thr Arg Lys
1      5      10      15
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Pro Tyr Thr Ile Thr Lys Gln Arg Glu Arg Trp Thr Glu Glu Glu His
20      25      30
aat aga ttc att gaa gct ttg agg ctt tat ggt aga gca tgg cag aag      381
Asn Arg Phe Ile Glu Ala Leu Arg Leu Tyr Gly Arg Ala Trp Gln Lys
35      40      45
att gaa gaa cat gta gca aca aaa act gct gtc cag ata aga agt cac      429
Ile Glu Glu His Val Ala Thr Lys Thr Ala Val Gln Ile Arg Ser His
50      55      60
gct cag aaa ttt ttc tcc aag gta gag aaa gag gct gaa gct aaa ggt      477
Ala Gln Lys Phe Phe Ser Lys Val Glu Lys Glu Ala Glu Ala Lys Gly
65      70      75      80
gta gct atg ggt caa gcg cta gac ata gct att cct cct cca cgg cct      525
Val Ala Met Gly Gln Ala Leu Asp Ile Ala Ile Pro Pro Pro Arg Pro
85      90      95
aag cgt aaa cca aac aat cct tat cct cga aag acg gga agt gga acg      573
Lys Arg Lys Pro Asn Asn Pro Tyr Pro Arg Lys Thr Gly Ser Gly Thr
100      105      110
atc ctt atg tca aaa acg ggt gtg aat gat gga aaa gag tcc ctt gga      621
Ile Leu Met Ser Lys Thr Gly Val Asn Asp Gly Lys Glu Ser Leu Gly
115      120      125
tca gaa aaa gtg tcg cat cct gag atg gcc aat gaa gat cga caa caa      669
Ser Glu Lys Val Ser His Pro Glu Met Ala Asn Glu Asp Arg Gln Gln
130      135      140
tca aag cct gaa gag aaa act ctg cag gaa gac aac tgt tca gat tgt      717
Ser Lys Pro Glu Glu Lys Thr Leu Gln Glu Asp Asn Cys Ser Asp Cys
145      150      155      160
ttc act cat cag tat ctc tct gct gca tcc tcc atg aat aaa agt tgt      765
Phe Thr His Gln Tyr Leu Ser Ala Ala Ser Ser Met Asn Lys Ser Cys
165      170      175
ata gag aca tca aac gca agc act ttc cgc gag ttc ttg cct tca cgg      813
Ile Glu Thr Ser Asn Ala Ser Thr Phe Arg Glu Phe Leu Pro Ser Arg
180      185      190
gaa gag gga agt cag aat aac agg gta aga aag gag tca aac tca gat      861
Glu Glu Gly Ser Gln Asn Asn Arg Val Arg Lys Glu Ser Asn Ser Asp
195      200      205
ttg aat gca aaa tct ctg gaa aac ggt aat gag caa gga cct cag act      909
Leu Asn Ala Lys Ser Leu Glu Asn Gly Asn Glu Gln Gly Pro Gln Thr
210      215      220
tat ccg atg cat atc cct gtg cta gtg cca ttg ggg agc tca ata aca      957
Tyr Pro Met His Ile Pro Val Leu Val Pro Leu Gly Ser Ser Ile Thr
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mbil9 Sequence Listing.ST25

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Val	Ala	Gly	Asp	Tyr	Gln	Ser	Phe	Pro	Asn	His	Ile	Met	Ser	Thr	Leu	
			260					265					270			
tta	caa	aca	ccg	gct	ctt	tat	act	gcc	gca	act	ttc	gcc	tca	tca	ttt	1101
Leu	Gln	Thr	Pro	Ala	Leu	Tyr	Thr	Ala	Ala	Thr	Phe	Ala	Ser	Ser	Phe	
			275				280					285				
tgg	cct	ccc	gat	tct	agt	ggg	ggc	tca	cct	ggt	cca	ggg	aac	tca	cct	1149
Trp	Pro	Pro	Asp	Ser	Ser	Gly	Gly	Ser	Pro	Val	Pro	Gly	Asn	Ser	Pro	
			290			295					300					
ccg	aat	ctg	gct	gcc	atg	gcc	gca	gcc	act	ggt	gca	gct	gct	agt	gct	1197
Pro	Asn	Leu	Ala	Ala	Met	Ala	Ala	Ala	Thr	Val	Ala	Ala	Ala	Ser	Ala	
305					310					315					320	
tgg	tgg	gct	gcc	aat	gga	tta	tta	cct	tta	tgt	gct	cct	ctt	agt	tca	1245
Trp	Trp	Ala	Ala	Asn	Gly	Leu	Leu	Pro	Leu	Cys	Ala	Pro	Leu	Ser	Ser	
				325				330						335		
ggg	ggg	ttc	act	agt	cat	cct	cca	tct	act	ttt	gga	cca	tca	tgt	gat	1293
Gly	Gly	Phe	Thr	Ser	His	Pro	Pro	Ser	Thr	Phe	Gly	Pro	Ser	Cys	Asp	
			340					345					350			
gta	gag	tac	aca	aaa	gca	agc	act	tta	caa	cat	ggg	tct	gtg	cag	agc	1341
Val	Glu	Tyr	Thr	Lys	Ala	Ser	Thr	Leu	Gln	His	Gly	Ser	Val	Gln	Ser	
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Arg	Glu	Gln	Glu	His	Ser	Gly	Ala	Ser	Lys	Ala	Arg	Ser	Ser	Leu	Asp	
			370			375					380					
tca	gag	gat	ggt	gaa	aat	aag	agt	aaa	cca	ggt	tgt	cat	gag	cag	cct	1437
Ser	Glu	Asp	Val	Glu	Asn	Lys	Ser	Lys	Pro	Val	Cys	His	Glu	Gln	Pro	
385					390					395					400	
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Ser	Ala	Thr	Pro	Glu	Ser	Asp	Ala	Lys	Gly	Ser	Asp	Gly	Ala	Gly	Asp	
				405					410					415		
aga	aaa	caa	ggt	gac	cgg	tcc	tcg	tgt	ggc	tca	aac	act	ccg	tcg	agt	1533
Arg	Lys	Gln	Val	Asp	Arg	Ser	Ser	Cys	Gly	Ser	Asn	Thr	Pro	Ser	Ser	
			420					425					430			
agt	gat	gat	ggt	gag	gcg	gat	gca	tca	gaa	agg	caa	gag	gat	ggc	acc	1581
Ser	Asp	Asp	Val	Glu	Ala	Asp	Ala	Ser	Glu	Arg	Gln	Glu	Asp	Gly	Thr	
			435				440					445				
aat	ggg	gag	gtg	aaa	gaa	acg	aat	gaa	gac	act	aat	aaa	cct	caa	act	1629
Asn	Gly	Glu	Val	Lys	Glu	Thr	Asn	Glu	Asp	Thr	Asn	Lys	Pro	Gln	Thr	
			450			455					460					
tca	gag	tcc	aat	gca	cgc	cgc	agt	aga	atc	agc	tcc	aat	ata	acc	gat	1677
Ser	Glu	Ser	Asn	Ala	Arg	Arg	Ser	Arg	Ile	Ser	Ser	Asn	Ile	Thr	Asp	
465					470					475					480	
cca	tgg	aag	tct	gtg	tct	gac	gag	ggg	cga	att	gcc	ttc	caa	gct	ctc	1725
Pro	Trp	Lys	Ser	Val	Ser	Asp	Glu	Gly	Arg	Ile	Ala	Phe	Gln	Ala	Leu	
				485				490						495		
ttc	tcc	aga	gag	gta	ttg	ccg	caa	agt	ttt	aca	tat	cga	gaa	gaa	cac	1773
Phe	Ser	Arg	Glu	Val	Leu	Pro	Gln	Ser	Phe	Thr	Tyr	Arg	Glu	Glu	His	
			500					505					510			
aga	gag	gaa	gaa	caa	caa	caa	caa	gaa	caa	aga	tat	cca	atg	gca	ctt	1821
Arg	Glu	Glu	Glu	Gln	Gln	Gln	Gln	Glu	Gln	Arg	Tyr	Pro	Met	Ala	Leu	
			515				520					525				
gat	ctt	aac	ttc	aca	gct	cag	tta	aca	cca	ggt	gat	gat	caa	gag	gag	1869
Asp	Leu	Asn	Phe	Thr	Ala	Gln	Leu	Thr	Pro	Val	Asp	Asp	Gln	Glu	Glu	
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mbi19 Sequence Listing.ST25

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Lys Arg Asn Thr Gly Phe Leu Gly Ile Gly Leu Asp Ala Ser Lys Leu 560
545 550 555

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Met Ser Arg Gly Arg Thr Gly Phe Lys Pro Tyr Lys Arg Cys Ser Met 575
565 570

gaa gcc aaa gaa agt aga atc ctc aac aac aat cct atc att cat gtg 2013
Glu Ala Lys Glu Ser Arg Ile Leu Asn Asn Asn Pro Ile Ile His Val 590
580 585

gaa cag aaa gat ccc aaa cgg atg cgg ttg gaa act caa gct tcc aca 2061
Glu Gln Lys Asp Pro Lys Arg Met Arg Leu Glu Thr Gln Ala Ser Thr 605
595 600

tga gactctatatt tcatctgata tgttgtttgt actctgtttt taagtatttca 2114

agaccactgc tacattttct ttttcttttg aggcctttgt atttgtttcc ttgtccatag 2174

tcttcctgta acatttgact ctgtattatt caacaaatca taaactgttt aatctttttt 2234

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Asn Arg Phe Ile Glu Ala Leu Arg Leu Tyr Gly Arg Ala Trp Gln Lys
35 40 45

Ile Glu Glu His Val Ala Thr Lys Thr Ala Val Gln Ile Arg Ser His
50 55 60

Ala Gln Lys Phe Phe Ser Lys Val Glu Lys Glu Ala Glu Ala Lys Gly
65 70 75 80

Val Ala Met Gly Gln Ala Leu Asp Ile Ala Ile Pro Pro Pro Arg Pro
85 90 95

Lys Arg Lys Pro Asn Asn Pro Tyr Pro Arg Lys Thr Gly Ser Gly Thr
100 105 110

Ile Leu Met Ser Lys Thr Gly Val Asn Asp Gly Lys Glu Ser Leu Gly
115 120 125

Ser Glu Lys Val Ser His Pro Glu Met Ala Asn Glu Asp Arg Gln Gln
130 135 140

Ser Lys Pro Glu Glu Lys Thr Leu Gln Glu Asp Asn Cys Ser Asp Cys
145 150 155 160

Phe Thr His Gln Tyr Leu Ser Ala Ala Ser Ser Met Asn Lys Ser Cys
165 170 175

mbi19 Sequence Listing.ST25

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 210 215 220
 Tyr Pro Met His Ile Pro Val Leu Val Pro Leu Gly Ser Ser Ile Thr
 225 230 235 240
 Ser Ser Leu Ser His Pro Pro Ser Glu Pro Asp Ser His Pro His Thr
 245 250 255
 Val Ala Gly Asp Tyr Gln Ser Phe Pro Asn His Ile Met Ser Thr Leu
 260 265 270
 Leu Gln Thr Pro Ala Leu Tyr Thr Ala Ala Thr Phe Ala Ser Ser Phe
 275 280 285
 Trp Pro Pro Asp Ser Ser Gly Gly Ser Pro Val Pro Gly Asn Ser Pro
 290 295 300
 Pro Asn Leu Ala Ala Met Ala Ala Ala Thr Val Ala Ala Ala Ser Ala
 305 310 315 320
 Trp Trp Ala Ala Asn Gly Leu Leu Pro Leu Cys Ala Pro Leu Ser Ser
 325 330 335
 Gly Gly Phe Thr Ser His Pro Pro Ser Thr Phe Gly Pro Ser Cys Asp
 340 345 350
 Val Glu Tyr Thr Lys Ala Ser Thr Leu Gln His Gly Ser Val Gln Ser
 355 360 365
 Arg Glu Gln Glu His Ser Glu Ala Ser Lys Ala Arg Ser Ser Leu Asp
 370 375 380
 Ser Glu Asp Val Glu Asn Lys Ser Lys Pro Val Cys His Glu Gln Pro
 385 390 395 400
 Ser Ala Thr Pro Glu Ser Asp Ala Lys Gly Ser Asp Gly Ala Gly Asp
 405 410 415
 Arg Lys Gln Val Asp Arg Ser Ser Cys Gly Ser Asn Thr Pro Ser Ser
 420 425 430
 Ser Asp Asp Val Glu Ala Asp Ala Ser Glu Arg Gln Glu Asp Gly Thr
 435 440 445
 Asn Gly Glu Val Lys Glu Thr Asn Glu Asp Thr Asn Lys Pro Gln Thr
 450 455 460
 Ser Glu Ser Asn Ala Arg Arg Ser Arg Ile Ser Ser Asn Ile Thr Asp
 465 470 475 480

mbil9 Sequence Listing.ST25

Pro Trp Lys Ser Val Ser Asp Glu Gly Arg Ile Ala Phe Gln Ala Leu
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Phe Ser Arg Glu Val Leu Pro Gln Ser Phe Thr Tyr Arg Glu Glu His
500 505 510

Arg Glu Glu Glu Gln Gln Gln Gln Glu Gln Arg Tyr Pro Met Ala Leu
515 520 525

Asp Leu Asn Phe Thr Ala Gln Leu Thr Pro Val Asp Asp Gln Glu Glu
530 535 540

Lys Arg Asn Thr Gly Phe Leu Gly Ile Gly Leu Asp Ala Ser Lys Leu
545 550 555 560

Met Ser Arg Gly Arg Thr Gly Phe Lys Pro Tyr Lys Arg Cys Ser Met
565 570 575

Glu Ala Lys Glu Ser Arg Ile Leu Asn Asn Asn Pro Ile Ile His Val
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Glu Gln Lys Asp Pro Lys Arg Met Arg Leu Glu Thr Gln Ala Ser Thr
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<223> G1930

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Met Asp Ala Met Ser Ser Val Asp Glu Ser Ser Thr
1 5 10
act aca gat tcc att ccg gcg aga aag tca tcg tct ccg gcg agt tta 159
Thr Thr Asp Ser Ile Pro Ala Arg Lys Ser Ser Ser Pro Ala Ser Leu
15 20 25
cta tat aga atg gga agc gga aca agc gtg gta ctt gat tca gag aac 207
Leu Tyr Arg Met Gly Ser Gly Thr Ser Val Val Leu Asp Ser Glu Asn
30 35 40
ggg gtc gaa gtc gaa gtc gaa gcc gaa tca aga aag ctt cct tct tca 255
Gly Val Glu Val Glu Val Glu Ala Glu Ser Arg Lys Leu Pro Ser Ser
45 50 55 60
aga ttc aaa ggt gtt gtt cct caa cca aat gga aga tgg gga gct cag 303
Arg Phe Lys Gly Val Val Pro Gln Pro Asn Gly Arg Trp Gly Ala Gln
65 70 75
att tac gag aaa cat caa cgc gtg tgg ctt ggt act ttc aac gag gaa 351
Ile Tyr Glu Lys His Gln Arg Val Trp Leu Gly Thr Phe Asn Glu Glu
80 85 90
gac gaa gca gct cgt gct tac gac gtc gcg gct cac cgt ttc cgt ggc 399
Asp Glu Ala Ala Arg Ala Tyr Asp Val Ala Ala His Arg Phe Arg Gly
95 100 105

mbi19 Sequence Listing.ST25

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aaa cac act tac aaa gaa gag tta gac caa agg aaa cgt aac cgt gac Lys His Thr Tyr Lys Glu Glu Leu Asp Gln Arg Lys Arg Asn Arg Asp 145 150 155	543
ggg aac gga aaa gag acg acg gcg ttt gct ttg gct tcg atg gtg gtt Gly Asn Gly Lys Glu Thr Thr Ala Phe Ala Leu Ala Ser Met Val Val 160 165 170	591
atg acg ggg ttt aaa acg gcg gag tta ctg ttt gag aaa acg gta acg Met Thr Gly Phe Lys Thr Ala Glu Leu Leu Phe Glu Lys Thr Val Thr 175 180 185	639
cca agt gac gtc ggg aaa cta aac cgt tta gtt ata cca aaa cac caa Pro Ser Asp Val Gly Lys Leu Asn Arg Leu Val Ile Pro Lys His Gln 190 195 200	687
gcg gag aaa cat ttt ccg tta ccg tta ggt aat aat aac gtc tcc gtt Ala Glu Lys His Phe Pro Leu Pro Leu Gly Asn Asn Asn Val Ser Val 205 210 215 220	735
aaa ggt atg ctg ttg aat ttc gaa gac gtt aac ggg aaa gtg tgg agg Lys Gly Met Leu Leu Asn Phe Glu Asp Val Asn Gly Lys Val Trp Arg 225 230 235	783
ttc cgt tac tct tat tgg aat agt agt caa agt tat gtg ttg acc aaa Phe Arg Tyr Ser Tyr Trp Asn Ser Ser Gln Ser Tyr Val Leu Thr Lys 240 245 250	831
ggg tgg agt aga ttc gtt aaa gag aag aga ctt tgt gct ggt gat ttg Gly Trp Ser Arg Phe Val Lys Glu Lys Arg Leu Cys Ala Gly Asp Leu 255 260 265	879
atc agt ttt aaa aga tcc aac gat caa gat caa aaa ttc ttt atc ggg Ile Ser Phe Lys Arg Ser Asn Asp Gln Asp Gln Lys Phe Phe Ile Gly 270 275 280	927
tgg aaa tcg aaa tcc ggg ttg gat cta gag acg ggt cgg gtt atg aga Trp Lys Ser Lys Ser Gly Leu Asp Leu Glu Thr Gly Arg Val Met Arg 285 290 295 300	975
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aca acg gag gtg tta atg tcg tcg tta agg tgt aag aag caa cga gtt Thr Thr Glu Val Leu Met Ser Ser Leu Arg Cys Lys Lys Gln Arg Val 320 325 330	1071
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mbil9 Sequence Listing.ST25

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Val Val Pro Gln Pro Asn Gly Arg Trp Gly Ala Gln Ile Tyr Glu Lys
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His Gln Arg Val Trp Leu Gly Thr Phe Asn Glu Glu Asp Glu Ala Ala
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Arg Ala Tyr Asp Val Ala Ala His Arg Phe Arg Gly Arg Asp Ala Val
      100             105             110

Thr Asn Phe Lys Asp Thr Thr Phe Glu Glu Glu Val Glu Phe Leu Asn
      115             120             125

Ala His Ser Lys Ser Glu Ile Val Asp Met Leu Arg Lys His Thr Tyr
      130             135             140

Lys Glu Glu Leu Asp Gln Arg Lys Arg Asn Arg Asp Gly Asn Gly Lys
      145             150             155             160

Glu Thr Thr Ala Phe Ala Leu Ala Ser Met Val Val Met Thr Gly Phe
      165             170             175

Lys Thr Ala Glu Leu Leu Phe Glu Lys Thr Val Thr Pro Ser Asp Val
      180             185             190

Gly Lys Leu Asn Arg Leu Val Ile Pro Lys His Gln Ala Glu Lys His
      195             200             205

Phe Pro Leu Pro Leu Gly Asn Asn Asn Val Ser Val Lys Gly Met Leu
      210             215             220

Leu Asn Phe Glu Asp Val Asn Gly Lys Val Trp Arg Phe Arg Tyr Ser
      225             230             235             240

Tyr Trp Asn Ser Ser Gln Ser Tyr Val Leu Thr Lys Gly Trp Ser Arg
      245             250             255

Phe Val Lys Glu Lys Arg Leu Cys Ala Gly Asp Leu Ile Ser Phe Lys
      260             265             270

Arg Ser Asn Asp Gln Asp Gln Lys Phe Phe Ile Gly Trp Lys Ser Lys
      275             280             285

Ser Gly Leu Asp Leu Glu Thr Gly Arg Val Met Arg Leu Phe Gly Val
      290             295             300

Asp Ile Ser Leu Asn Ala Val Val Val Val Lys Glu Thr Thr Glu Val
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mbil9 Sequence Listing.ST25

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 Met Asp Ser Ser Cys Ile Asp Glu Ile Ser Ser
 1 5 10
 tcc act tca gaa tct ttc tcc gcc acc acc gcc aag aag ctc tct cct 161
 Ser Thr Ser Glu Ser Phe Ser Ala Thr Thr Ala Lys Lys Leu Ser Pro
 15 20 25
 cct ccc gcg gcg gcg tta cgc ctc tac cgg atg gga agc ggc ggg agc 209
 Pro Pro Ala Ala Ala Leu Arg Leu Tyr Arg Met Gly Ser Gly Gly Ser
 30 35 40
 agc gtc gtg ttg gat ccc gag aac ggc cta gag acg gag tca cga aag 257
 Ser Val Val Leu Asp Pro Glu Asn Gly Leu Glu Thr Glu Ser Arg Lys
 45 50 55
 cta cca tct tca aaa tac aaa ggt gtt gtt cct cag cct aac gga aga 305
 Leu Pro Ser Ser Lys Tyr Lys Gly Val Val Pro Gln Pro Asn Gly Arg
 60 65 70 75
 tgg gga gct cag atc tac gag aag cac caa cga gta tgg ctc ggg act 353
 Trp Gly Ala Gln Ile Tyr Glu Lys His Gln Arg Val Trp Leu Gly Thr
 80 85 90
 ttc aac gag caa gaa gaa gct gct cgt tcc tac gac atc gca gct tgt 401
 Phe Asn Glu Gln Glu Glu Ala Ala Arg Ser Tyr Asp Ile Ala Ala Cys
 95 100 105
 aga ttc cgt ggc cgc gac gcc gtc gtc aac ttc aag aac gtt ctg gaa 449
 Arg Phe Arg Gly Arg Asp Ala Val Val Asn Phe Lys Asn Val Leu Glu
 110 115 120
 gac ggc gat tta gct ttt ctt gaa gct cac tca aag gcc gag atc gtc 497
 Asp Gly Asp Leu Ala Phe Leu Glu Ala His Ser Lys Ala Glu Ile Val
 125 130 135
 gac atg ttg aga aaa cac act tac gcc gac gag ctt gaa cag aac aat 545
 Asp Met Leu Arg Lys His Thr Tyr Ala Asp Glu Leu Glu Gln Asn Asn
 140 145 150 155
 aaa cgg cag ttg ttt ctc tcc gtc gac gct aac gga aaa cgt aac gga 593
 Lys Arg Gln Leu Phe Leu Ser Val Asp Ala Asn Gly Lys Arg Asn Gly
 160 165 170
 tcg agt act act caa aac gac aaa gtt tta aag acg tgt gaa gtt ctt 641
 Ser Ser Thr Thr Gln Asn Asp Lys Val Leu Lys Thr Cys Glu Val Leu
 175 180 185
 ttc gag aag gct gtt aca cct agc gac gtt ggg aag cta aac cgt ctc 689
 Phe Glu Lys Ala Val Thr Pro Ser Asp Val Gly Lys Leu Asn Arg Leu
 190 195 200
 gtg ata cct aaa caa cac gcc gag aaa cac ttt ccg tta ccg tca ccg 737
 Val Ile Pro Lys Gln His Ala Glu Lys His Phe Pro Leu Pro Ser Pro
 205 210 215

mbi19 Sequence Listing.ST25

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Ser Pro Ala Val Thr Lys Gly Val Leu Ile Asn Phe Glu Asp Val Asn	
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Gly Lys Val Trp Arg Phe Arg Tyr Ser Tyr Trp Asn Ser Ser Gln Ser	
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tac gtg ttg acc aag gga tgg agt cga ttc gtc aag gag aag aat ctt	881
Tyr Val Leu Thr Lys Gly Trp Ser Arg Phe Val Lys Glu Lys Asn Leu	
255 260 265	
cga gcc ggt gat gtt gtt act ttc gag aga tcg acc gga cta gag cgg	929
Arg Ala Gly Asp Val Val Thr Phe Glu Arg Ser Thr Gly Leu Glu Arg	
270 275 280	
cag tta tat att gat tgg aaa gtt cgg tct ggt ccg aga gaa aac ccg	977
Gln Leu Tyr Ile Asp Trp Lys Val Arg Ser Gly Pro Arg Glu Asn Pro	
285 290 295	
ggt cag gtg gtg gtt cgg ctt ttc gga gtt gat atc ttt aat gtg acc	1025
Val Gln Val Val Val Arg Leu Phe Gly Val Asp Ile Phe Asn Val Thr	
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acc gtg aag cca aac gac gtc gtg gcc gtt tgc ggt gga aag aga tct	1073
Thr Val Lys Pro Asn Asp Val Val Ala Val Cys Gly Gly Lys Arg Ser	
320 325 330	
cga gat gtt gat gat atg ttt gcg tta cgg tgt tcc aag aag cag gcg	1121
Arg Asp Val Asp Asp Met Phe Ala Leu Arg Cys Ser Lys Lys Gln Ala	
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Ile Ile Asn Ala Leu	
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Pro Glu Asn Gly Leu Glu Thr Glu Ser Arg Lys Leu Pro Ser Ser Lys	
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Tyr Lys Gly Val Val Pro Gln Pro Asn Gly Arg Trp Gly Ala Gln Ile	
65 70 75 80	
Tyr Glu Lys His Gln Arg Val Trp Leu Gly Thr Phe Asn Glu Gln Glu	
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Glu Ala Ala Arg Ser Tyr Asp Ile Ala Ala Cys Arg Phe Arg Gly Arg	
100 105 110	

mbi19 Sequence Listing.ST25

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 130 135 140
 His Thr Tyr Ala Asp Glu Leu Glu Gln Asn Asn Lys Arg Gln Leu Phe
 145 150 155 160
 Leu Ser Val Asp Ala Asn Gly Lys Arg Asn Gly Ser Ser Thr Thr Gln
 165 170 175
 Asn Asp Lys Val Leu Lys Thr Cys Glu Val Leu Phe Glu Lys Ala Val
 180 185 190
 Thr Pro Ser Asp Val Gly Lys Leu Asn Arg Leu Val Ile Pro Lys Gln
 195 200 205
 His Ala Glu Lys His Phe Pro Leu Pro Ser Pro Ser Pro Ala Val Thr
 210 215 220
 Lys Gly Val Leu Ile Asn Phe Glu Asp Val Asn Gly Lys Val Trp Arg
 225 230 235 240
 Phe Arg Tyr Ser Tyr Trp Asn Ser Ser Gln Ser Tyr Val Leu Thr Lys
 245 250 255
 Gly Trp Ser Arg Phe Val Lys Glu Lys Asn Leu Arg Ala Gly Asp Val
 260 265 270
 Val Thr Phe Glu Arg Ser Thr Gly Leu Glu Arg Gln Leu Tyr Ile Asp
 275 280 285
 Trp Lys Val Arg Ser Gly Pro Arg Glu Asn Pro Val Gln Val Val Val
 290 295 300
 Arg Leu Phe Gly Val Asp Ile Phe Asn Val Thr Thr Val Lys Pro Asn
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 <223> G993

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mbi19 Sequence Listing.ST25

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Leu Ser Ser Pro Pro Ala Thr Ser Met Arg Leu Tyr Arg Met Gly Ser				
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ggc gga agc agc gtc gtt ttg gat tca gag aac ggc gtc gag acc gag	194			
Gly Gly Ser Ser Val Val Leu Asp Ser Glu Asn Gly Val Glu Thr Glu				
50 55 60				
tca cgt aag ctt cct tcg tcg aaa tat aaa ggc gtt gtg cct cag cct	242			
Ser Arg Lys Leu Pro Ser Ser Lys Tyr Lys Gly Val Val Pro Gln Pro				
65 70 75				
aac gga aga tgg gga gct cag att tac gag aag cat cag cga gtt tgg	290			
Asn Gly Arg Trp Gly Ala Gln Ile Tyr Glu Lys His Gln Arg Val Trp				
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ctc ggt act ttc aac gag gaa gaa gaa gct gcg tct tct tac gac atc	338			
Leu Gly Thr Phe Asn Glu Glu Glu Glu Ala Ala Ser Ser Tyr Asp Ile				
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gcc gtg agg aga ttc cgc gcc cgc gac gcc gtc act aac ttc aaa tct	386			
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Gln Val Asp Gly Asn Asp Ala Glu Ser Ala Phe Leu Asp Ala His Ser				
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Lys Ala Glu Ile Val Asp Met Leu Arg Lys His Thr Tyr Ala Asp Glu				
145 150 155				
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Phe Glu Gln Ser Arg Arg Lys Phe Val Asn Gly Asp Gly Lys Arg Ser				
160 165 170 175				
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Gly Leu Glu Thr Ala Thr Tyr Gly Asn Asp Ala Val Leu Arg Ala Arg				
180 185 190				
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Glu Val Leu Phe Glu Lys Thr Val Thr Pro Ser Asp Val Gly Lys Leu				
195 200 205				
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Asn Arg Leu Val Ile Pro Lys Gln His Ala Glu Lys His Phe Pro Leu				
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225 230 235				
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Val Leu Ile Asn Leu Glu Asp Arg Thr Gly Lys Val Trp Arg Phe Arg				
240 245 250 255				
tac agt tac tgg aac agc agt caa agt tac gtg ttg acc aag ggc tgg	818			
Tyr Ser Tyr Trp Asn Ser Ser Gln Ser Tyr Val Leu Thr Lys Gly Trp				
260 265 270				
agc cgg ttc gtt aaa gag aag aat ctt cga gcc ggt gat gtg gtt tgt	866			
Ser Arg Phe Val Lys Glu Lys Asn Leu Arg Ala Gly Asp Val Val Cys				
275 280 285				
ttc gag aga tca acc gga cca gac ccg caa ttg tat atc cac tgg aaa	914			
Phe Glu Arg Ser Thr Gly Pro Asp Arg Gln Leu Tyr Ile His Trp Lys				
290 295 300				
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mbi19 Sequence Listing.ST25

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Ile	Phe	Asn	Val	Ser	Asn	Glu	Lys	Pro	Asn	Asp	Val	Ala	Val	Glu	Cys		
320					325					330					335		
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Val	Gly	Lys	Lys	Arg	Ser	Arg	Glu	Asp	Asp	Leu	Phe	Ser	Leu	Gly	Cys		
				340					345					350			
tcc	aag	aag	cag	gcg	att	atc	aac	atc	ttg	tga	caaattcttt	ttttttgggt	1111				
Ser	Lys	Lys	Gln	Ala	Ile	Ile	Asn	Ile	Leu								
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aaaaaaaaa						1239											

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 <213> Arabidopsis thaliana

<400> 58

Met	Glu	Tyr	Ser	Cys	Val	Asp	Asp	Ser	Ser	Thr	Thr	Ser	Glu	Ser	Leu		
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Ser	Ile	Ser	Thr	Thr	Pro	Lys	Pro	Thr	Thr	Thr	Thr	Glu	Lys	Lys	Leu		
			20					25					30				
Ser	Ser	Pro	Pro	Ala	Thr	Ser	Met	Arg	Leu	Tyr	Arg	Met	Gly	Ser	Gly		
		35					40					45					
Gly	Ser	Ser	Val	Val	Leu	Asp	Ser	Glu	Asn	Gly	Val	Glu	Thr	Glu	Ser		
	50					55					60						
Arg	Lys	Leu	Pro	Ser	Ser	Lys	Tyr	Lys	Gly	Val	Val	Pro	Gln	Pro	Asn		
65					70					75					80		
Gly	Arg	Trp	Gly	Ala	Gln	Ile	Tyr	Glu	Lys	His	Gln	Arg	Val	Trp	Leu		
				85					90					95			
Gly	Thr	Phe	Asn	Glu	Glu	Glu	Glu	Ala	Ala	Ser	Ser	Tyr	Asp	Ile	Ala		
			100					105					110				
Val	Arg	Arg	Phe	Arg	Gly	Arg	Asp	Ala	Val	Thr	Asn	Phe	Lys	Ser	Gln		
			115				120					125					
Val	Asp	Gly	Asn	Asp	Ala	Glu	Ser	Ala	Phe	Leu	Asp	Ala	His	Ser	Lys		
	130					135					140						
Ala	Glu	Ile	Val	Asp	Met	Leu	Arg	Lys	His	Thr	Tyr	Ala	Asp	Glu	Phe		
145					150					155					160		
Glu	Gln	Ser	Arg	Arg	Lys	Phe	Val	Asn	Gly	Asp	Gly	Lys	Arg	Ser	Gly		
				165					170					175			
Leu	Glu	Thr	Ala	Thr	Tyr	Gly	Asn	Asp	Ala	Val	Leu	Arg	Ala	Arg	Glu		
			180					185					190				

mbil9 Sequence Listing.ST25

Val Leu Phe Glu Lys Thr Val Thr Pro Ser Asp Val Gly Lys Leu Asn
195 200 205

Arg Leu Val Ile Pro Lys Gln His Ala Glu Lys His Phe Pro Leu Pro
210 215 220

Ala Met Thr Thr Ala Met Gly Met Asn Pro Ser Pro Thr Lys Gly Val
225 230 235 240

Leu Ile Asn Leu Glu Asp Arg Thr Gly Lys Val Trp Arg Phe Arg Tyr
245 250 255

Ser Tyr Trp Asn Ser Ser Gln Ser Tyr Val Leu Thr Lys Gly Trp Ser
260 265 270

Arg Phe Val Lys Glu Lys Asn Leu Arg Ala Gly Asp Val Val Cys Phe
275 280 285

Glu Arg Ser Thr Gly Pro Asp Arg Gln Leu Tyr Ile His Trp Lys Val
290 295 300

Arg Ser Ser Pro Val Gln Thr Val Val Arg Leu Phe Gly Val Asn Ile
305 310 315 320

Phe Asn Val Ser Asn Glu Lys Pro Asn Asp Val Ala Val Glu Cys Val
325 330 335

Gly Lys Lys Arg Ser Arg Glu Asp Asp Leu Phe Ser Leu Gly Cys Ser
340 345 350

Lys Lys Gln Ala Ile Ile Asn Ile Leu
355 360

<210> 59
<211> 803
<212> DNA
<213> Arabidopsis thaliana

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<222> (35)..(658)
<223> G41

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Ser Pro Val Ser Ser Gly Gly Asp Tyr Ser Pro Lys Leu Ala Thr Ser
10 15 20

tgc ccc aag aaa cca gcg gga agg aag aag ttt cgt gag act cgt cac 151
Cys Pro Lys Lys Pro Ala Gly Arg Lys Lys Phe Arg Glu Thr Arg His
25 30 35

cca att tac aga gga gtt cgt caa aga aac tcc ggt aag tgg gtg tgt 199
Pro Ile Tyr Arg Gly Val Arg Gln Arg Asn Ser Gly Lys Trp Val Cys
40 45 50 55

gag ttg aga gag cca aac aag aaa acg agg att tgg ctc ggg act ttc 247

mbil9 Sequence Listing.ST25

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 60 65 70
 caa acc gct gag atg gca gct cgt gct cac gac gtc gcc gcc ata gct 295
 Gln Thr Ala Glu Met Ala Ala Arg Ala His Asp Val Ala Ala Ile Ala
 75 80 85
 ctc cgt ggc aga tct gcc tgt ctc aat ttc gct gac tcg gct tgg cgg 343
 Leu Arg Gly Arg Ser Ala Cys Leu Asn Phe Ala Asp Ser Ala Trp Arg
 90 95 100
 cta cga atc ccg gaa tca acc tgt gcc aag gaa atc caa aag gcg gcg 391
 Leu Arg Ile Pro Glu Ser Thr Cys Ala Lys Glu Ile Gln Lys Ala Ala
 105 110 115
 gct gaa gcc gcg ttg aat ttt caa gat gag atg tgt cat atg acg acg 439
 Ala Glu Ala Ala Leu Asn Phe Gln Asp Glu Met Cys His Met Thr Thr
 120 125 130 135
 gat gct cat ggt ctt gac atg gag gag acc ttg gtg gag gct att tat 487
 Asp Ala His Gly Leu Asp Met Glu Glu Thr Leu Val Glu Ala Ile Tyr
 140 145 150
 acg ccg gaa cag agc caa gat gcg ttt tat atg gat gaa gag gcg atg 535
 Thr Pro Glu Gln Ser Gln Asp Ala Phe Tyr Met Asp Glu Glu Ala Met
 155 160 165
 ttg ggg atg tct agt ttg ttg gat aac atg gcc gaa ggg atg ctt tta 583
 Leu Gly Met Ser Ser Leu Leu Asp Asn Met Ala Glu Gly Met Leu Leu
 170 175 180
 ccg tcg ccg tcg gtt caa tgg aac tat aat ttt gat gtc gag gga gat 631
 Pro Ser Pro Ser Val Gln Trp Asn Tyr Asn Phe Asp Val Glu Gly Asp
 185 190 195
 gat gac gtg tcc tta tgg agc tat taa aattcgattt ttatttccat 678
 Asp Asp Val Ser Leu Trp Ser Tyr
 200 205
 ttttggtatt atagcttttt atacatttga tcctttttta gaatggatct tcttcttttt 738
 ttggttgtga gaaacgaatg taaatggtaa aagttgttgt caaatgcaaa tgtttttgag 798
 tgcag 803

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 <211> 207
 <212> PRT
 <213> Arabidopsis thaliana

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 20 25 30
 Lys Phe Arg Glu Thr Arg His Pro Ile Tyr Arg Gly Val Arg Gln Arg
 35 40 45
 Asn Ser Gly Lys Trp Val Cys Glu Leu Arg Glu Pro Asn Lys Lys Thr
 50 55 60
 Arg Ile Trp Leu Gly Thr Phe Gln Thr Ala Glu Met Ala Ala Arg Ala
 65 70 75 80
 His Asp Val Ala Ala Ile Ala Leu Arg Gly Arg Ser Ala Cys Leu Asn
 85 90 95

mbi19 Sequence Listing.ST25

Phe Ala Asp Ser Ala Trp Arg Leu Arg Ile Pro Glu Ser Thr Cys Ala
 100 105 110

Lys Glu Ile Gln Lys Ala Ala Ala Glu Ala Ala Leu Asn Phe Gln Asp
 115 120 125

Glu Met Cys His Met Thr Thr Asp Ala His Gly Leu Asp Met Glu Glu
 130 135 140

Thr Leu Val Glu Ala Ile Tyr Thr Pro Glu Gln Ser Gln Asp Ala Phe
 145 150 155 160

Tyr Met Asp Glu Glu Ala Met Leu Gly Met Ser Ser Leu Leu Asp Asn
 165 170 175

Met Ala Glu Gly Met Leu Leu Pro Ser Pro Ser Val Gln Trp Asn Tyr
 180 185 190

Asn Phe Asp Val Glu Gly Asp Asp Asp Val Ser Leu Trp Ser Tyr
 195 200 205

<210> 61
 <211> 929
 <212> DNA
 <213> Arabidopsis thaliana

<220>
 <221> CDS
 <222> (164)..(805)
 <223> G40

<400> 61
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 agacagatat actatctttt attaatccaa aaagactgag aactctagta actacgtact 120
 acttaaacct tatccagttt cttgaaacag agtactctga tca atg aac tca ttt 175
 Met Asn Ser Phe
 1
 tca gct ttt tct gaa atg ttt ggc tcc gat tac gag cct caa ggc gga 223
 Ser Ala Phe Ser Glu Met Phe Gly Ser Asp Tyr Glu Pro Gln Gly Gly
 5 10 15 20
 gat tat tgt ccg acg ttg gcc acg agt tgt ccg aag aaa ccg gcg ggc 271
 Asp Tyr Cys Pro Thr Leu Ala Thr Ser Cys Pro Lys Lys Pro Ala Gly
 25 30 35
 cgt aag aag ttt cgt gag act cgt cac cca att tac aga gga gtt cgt 319
 Arg Lys Lys Phe Arg Glu Thr Arg His Pro Ile Tyr Arg Gly Val Arg
 40 45 50
 caa aga aac tcc ggt aag tgg gtt tct gaa gtg aga gag cca aac aag 367
 Gln Arg Asn Ser Gly Lys Trp Val Ser Glu Val Arg Glu Pro Asn Lys
 55 60 65
 aaa acc agg att tgg ctc ggg act ttc caa acc gct gag atg gca gct 415
 Lys Thr Arg Ile Trp Leu Gly Thr Phe Gln Thr Ala Glu Met Ala Ala
 70 75 80
 cgt gct cac gac gtc gct gca tta gcc ctc cgt ggc cga tca gca tgt 463
 Arg Ala His Asp Val Ala Ala Leu Ala Leu Arg Gly Arg Ser Ala Cys
 85 90 95 100
 ctc aac ttc gct gac tcg gct tgg cgg cta cga atc ccg gag tca aca 511

mbi19 Sequence Listing.ST25

Leu Asn Phe Ala Asp Ser Ala Trp Arg Leu Arg Ile Pro Glu Ser Thr
 105 110 115

tgc gcc aag gat atc caa aaa gcg gct gct gaa gcg gcg ttg gct ttt 559
 Cys Ala Lys Asp Ile Gln Lys Ala Ala Glu Ala Ala Leu Ala Phe
 120 125 130

caa gat gag acg tgt gat acg acg acc acg aat cat ggc ctg gac atg 607
 Gln Asp Glu Thr Cys Asp Thr Thr Thr Asn His Gly Leu Asp Met
 135 140 145

gag gag acg atg gtg gaa gct att tat aca ccg gaa cag agc gaa ggt 655
 Glu Glu Thr Met Val Glu Ala Ile Tyr Thr Pro Glu Gln Ser Glu Gly
 150 155 160

gcg ttt tat atg gat gag gag aca atg ttt ggg atg ccg act ttg ttg 703
 Ala Phe Tyr Met Asp Glu Glu Thr Met Phe Gly Met Pro Thr Leu Leu
 165 170 175 180

gat aat atg gct gaa ggc atg ctt tta ccg ccg ccg tct gtt caa tgg 751
 Asp Asn Met Ala Glu Gly Met Leu Leu Pro Pro Pro Ser Val Gln Trp
 185 190 195

aat cat aat tat gac ggc gaa gga gat ggt gac gtg tcg ctt tgg agt 799
 Asn His Asn Tyr Asp Gly Glu Gly Asp Gly Asp Val Ser Leu Trp Ser
 200 205 210

tac taa tattcgatag tcgtttccat ttttgtacta tagtttgaaa atattctagt 855
 Tyr

tcctttttttt agaatgggttc cttcatttta ttttatttta ttgttgtaga aacgagtgga 915

aaataattca atac 929

<210> 62
 <211> 213
 <212> PRT
 <213> Arabidopsis thaliana

<400> '62

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Pro Gln Gly Gly Asp Tyr Cys Pro Thr Leu Ala Thr Ser Cys Pro Lys
 20 25 30

Lys Pro Ala Gly Arg Lys Lys Phe Arg Glu Thr Arg His Pro Ile Tyr
 35 40 45

Arg Gly Val Arg Gln Arg Asn Ser Gly Lys Trp Val Ser Glu Val Arg
 50 55 60

Glu Pro Asn Lys Lys Thr Arg Ile Trp Leu Gly Thr Phe Gln Thr Ala
 65 70 75 80

Glu Met Ala Ala Arg Ala His Asp Val Ala Ala Leu Ala Leu Arg Gly
 85 90 95

Arg Ser Ala Cys Leu Asn Phe Ala Asp Ser Ala Trp Arg Leu Arg Ile
 100 105 110

Pro Glu Ser Thr Cys Ala Lys Asp Ile Gln Lys Ala Ala Ala Glu Ala
 115 120 125

mbil9 Sequence Listing.ST25

Ala Leu Ala Phe Gln Asp Glu Thr Cys Asp Thr Thr Thr Asn His
 130 135 140

Gly Leu Asp Met Glu Glu Thr Met Val Glu Ala Ile Tyr Thr Pro Glu
 145 150 155 160

Gln Ser Glu Gly Ala Phe Tyr Met Asp Glu Glu Thr Met Phe Gly Met
 165 170 175

Pro Thr Leu Leu Asp Asn Met Ala Glu Gly Met Leu Leu Pro Pro Pro
 180 185 190

Ser Val Gln Trp Asn His Asn Tyr Asp Gly Glu Gly Asp Gly Asp Val
 195 200 205

Ser Leu Trp Ser Tyr
 210

<210> 63
 <211> 908
 <212> DNA
 <213> Arabidopsis thaliana

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 <222> (119)..(769)
 <223> G42

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 atg aac tca ttt tct gct ttt tct gaa atg ttt ggc tcc gat tac gag 166
 Met Asn Ser Phe Ser Ala Phe Ser Glu Met Phe Gly Ser Asp Tyr Glu
 1 5 10 15
 tct tcg gtt tcc tca ggc ggt gat tat att ccg acg ctt gcg agc agc 214
 Ser Ser Val Ser Ser Gly Gly Asp Tyr Ile Pro Thr Leu Ala Ser Ser
 20 25 30
 tgc ccc aag aaa ccg gcg ggt cgt aag aag ttt cgt gag act cgt cac 262
 Cys Pro Lys Lys Pro Ala Gly Arg Lys Lys Phe Arg Glu Thr Arg His
 35 40 45
 cca ata tac aga gga gtt cgt cgg aga aac tcc ggt aag tgg gtt tgt 310
 Pro Ile Tyr Arg Gly Val Arg Arg Arg Asn Ser Gly Lys Trp Val Cys
 50 55 60
 gag gtt aga gaa cca aac aag aaa aca agg att tgg ctc gga aca ttt 358
 Glu Val Arg Glu Pro Asn Lys Lys Thr Arg Ile Trp Leu Gly Thr Phe
 65 70 75 80
 caa acc gct gag atg gca gct cga gct cac gac gtt gcc gct tta gcc 406
 Gln Thr Ala Glu Met Ala Ala Arg Ala His Asp Val Ala Ala Leu Ala
 85 90 95
 ctt cgt ggc cga tca gcc tgt ctc aat ttc gct gac tcg gct tgg aga 454
 Leu Arg Gly Arg Ser Ala Cys Leu Asn Phe Ala Asp Ser Ala Trp Arg
 100 105 110
 ctc cga atc ccg gaa tca act tgc gct aag gac atc caa aag gcg gcg 502
 Leu Arg Ile Pro Glu Ser Thr Cys Ala Lys Asp Ile Gln Lys Ala Ala
 115 120 125
 gct gaa gct gcg ttg gcg ttt cag gat gag atg tgt gat gcg acg acg 550
 Ala Glu Ala Ala Leu Ala Phe Gln Asp Glu Met Cys Asp Ala Thr Thr
 130 135 140

mbi19 Sequence Listing.ST25

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gat cat ggc ttc gac atg gag gag acg ttg gtg gag gct att tac acg      598
Asp His Gly Phe Asp Met Glu Glu Thr Leu Val Glu Ala Ile Tyr Thr
145                      150                      155                      160

gcg gaa cag agc gaa aat gcg ttt tat atg cac gat gag gcg atg ttt      646
Ala Glu Gln Ser Glu Asn Ala Phe Tyr Met His Asp Glu Ala Met Phe
                      165                      170                      175

gag atg ccg agt ttg ttg gct aat atg gca gaa ggg atg ctt ttg ccg      694
Glu Met Pro Ser Leu Leu Ala Asn Met Ala Glu Gly Met Leu Leu Pro
                      180                      185                      190

ctt ccg tcc gta cag tgg aat cat aat cat gaa gtc gac ggc gat gat      742
Leu Pro Ser Val Gln Trp Asn His Asn His Glu Val Asp Gly Asp Asp
                      195                      200                      205

gac gac gta tcg tta tgg agt tat taa aactcagatt attatttcca      789
Asp Asp Val Ser Leu Trp Ser Tyr
210                      215

tttttagtac gatacttttt attttattat tattttttaga tcctttttta gaatggaatc      849

tncattatgt ttgtaaaact gagaaacgag tgtaaattaa attgattcag tttcagtat      908

<210> 64
<211> 216
<212> PRT
<213> Arabidopsis thaliana

<400> 64

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Ser Ser Val Ser Ser Gly Gly Asp Tyr Ile Pro Thr Leu Ala Ser Ser
20                      25                      30

Cys Pro Lys Lys Pro Ala Gly Arg Lys Lys Phe Arg Glu Thr Arg His
35                      40                      45

Pro Ile Tyr Arg Gly Val Arg Arg Arg Asn Ser Gly Lys Trp Val Cys
50                      55                      60

Glu Val Arg Glu Pro Asn Lys Lys Thr Arg Ile Trp Leu Gly Thr Phe
65                      70                      75                      80

Gln Thr Ala Glu Met Ala Ala Arg Ala His Asp Val Ala Ala Leu Ala
85                      90                      95

Leu Arg Gly Arg Ser Ala Cys Leu Asn Phe Ala Asp Ser Ala Trp Arg
100                     105                     110

Leu Arg Ile Pro Glu Ser Thr Cys Ala Lys Asp Ile Gln Lys Ala Ala
115                     120                     125

Ala Glu Ala Ala Leu Ala Phe Gln Asp Glu Met Cys Asp Ala Thr Thr
130                     135                     140

Asp His Gly Phe Asp Met Glu Glu Thr Leu Val Glu Ala Ile Tyr Thr
145                     150                     155                     160

Ala Glu Gln Ser Glu Asn Ala Phe Tyr Met His Asp Glu Ala Met Phe
165                     170                     175

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mbil9 Sequence Listing.ST25

Glu Met Pro Ser Leu Leu Ala Asn Met Ala Glu Gly Met Leu Leu Pro
 180 185 190

Leu Pro Ser Val Gln Trp Asn His Asn His Glu Val Asp Gly Asp Asp
 195 200 205

Asp Asp Val Ser Leu Trp Ser Tyr
 210 215

<210> 65
 <211> 1407
 <212> DNA
 <213> Arabidopsis thaliana

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 <222> (191)..(1351)
 <223> G1127

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 cgagaattaa gccgaaagaa acaatctttg agtttgattt cttcttcctt ccttctctct 180
 ctctgctcta atg gat tcc aga gac atc cca ccg tca cat aac cag ctt 229
 Met Asp Ser Arg Asp Ile Pro Pro Ser His Asn Gln Leu
 1 5 10
 caa cca cca ccg gga atg tta atg tct cat tac cgt aac cct aac gcc 277
 Gln Pro Pro Pro Gly Met Leu Met Ser His Tyr Arg Asn Pro Asn Ala
 15 20 25
 gcc gct tca cca tta atg gtt ccc act tcc aca tct caa ccg att caa 325
 Ala Ala Ser Pro Leu Met Val Pro Thr Ser Thr Ser Gln Pro Ile Gln
 30 35 40 45
 cac cct cgt ctt cct ttt ggc aat caa caa caa tct caa acg ttt cat 373
 His Pro Arg Leu Pro Phe Gly Asn Gln Gln Gln Ser Gln Thr Phe His
 50 55 60
 cag cag caa caa caa caa atg gat cag aag act ctt gaa tct ctt gga 421
 Gln Gln Gln Gln Gln Gln Met Asp Gln Lys Thr Leu Glu Ser Leu Gly
 65 70 75
 ttt ggt gat gga tca cct tct tct caa ccg atg cga ttc ggg atc gat 469
 Phe Gly Asp Gly Ser Pro Ser Ser Gln Pro Met Arg Phe Gly Ile Asp
 80 85 90
 gat cag aat cag caa ctg caa gtg aag aag aag cga gga agg ccg aga 517
 Asp Gln Asn Gln Gln Leu Gln Val Lys Lys Lys Arg Gly Arg Pro Arg
 95 100 105
 aag tat act cct gat ggt agc att gct tta ggt tta gct cct acg tct 565
 Lys Tyr Thr Pro Asp Gly Ser Ile Ala Leu Gly Leu Ala Pro Thr Ser
 110 115 120 125
 cct ctt ctc tct gca gct tct aat tct tac ggt gag ggt ggt gtt gga 613
 Pro Leu Leu Ser Ala Ala Ser Asn Ser Tyr Gly Glu Gly Gly Val Gly
 130 135 140
 gat agt ggt gga aat gga aac tct gtt gat cca cct gtt aaa cgt aac 661
 Asp Ser Gly Gly Asn Gly Asn Ser Val Asp Pro Pro Val Lys Arg Asn
 145 150 155
 aga gga agg cct cct ggt tct agt aag aaa cag ctt gat gct tta gga 709
 Arg Gly Arg Pro Pro Gly Ser Ser Lys Lys Gln Leu Asp Ala Leu Gly
 160 165 170

mbil9 Sequence Listing.ST25

gga act tca gga gtt ggg ttt aca cct cat gtc att gaa gtg aac aca Gly Thr Ser Gly Val Gly Phe Thr Pro His Val Ile Glu Val Asn Thr 175 180 185	757
gga gag gac ata gcg tca aag gtg atg gct ttt tcg gat caa ggg tca Gly Glu Asp Ile Ala Ser Lys Val Met Ala Phe Ser Asp Gln Gly Ser 190 195 200 205	805
aga aca att tgt att ctc tct gca agt ggt gca gtt tct aga gtg atg Arg Thr Ile Cys Ile Leu Ser Ala Ser Gly Ala Val Ser Arg Val Met 210 215 220	853
ctt cgt caa gct tct cat tct agt gga atc gtt act tat gag gga cga Leu Arg Gln Ala Ser His Ser Ser Gly Ile Val Thr Tyr Glu Gly Arg 225 230 235	901
ttt gag atc att act ctc tca ggc tca gtc ttg aat tat gag gta aat Phe Glu Ile Ile Thr Leu Ser Gly Ser Val Leu Asn Tyr Glu Val Asn 240 245 250	949
ggt tcc acc aac aga agt ggt aac ttg agt gtg gct ttg gct gga cct Gly Ser Thr Asn Arg Ser Gly Asn Leu Ser Val Ala Leu Ala Gly Pro 255 260 265	997
gat ggc ggc atc gta ggt ggc agt gta gtt ggt aat cta gta gct gca Asp Gly Gly Ile Val Gly Gly Ser Val Val Gly Asn Leu Val Ala Ala 270 275 280 285	1045
aca caa gtc cag gtg ata gtg gga agc ttt gtt gca gaa gca aag aaa Thr Gln Val Gln Val Ile Val Gly Ser Phe Val Ala Glu Ala Lys Lys 290 295 300	1093
ccg aaa caa agt agt gtt aac att gct cgg ggg cag aat cct gaa ccg Pro Lys Gln Ser Ser Val Asn Ile Ala Arg Gly Gln Asn Pro Glu Pro 305 310 315	1141
gct tca gcg ccg gct aac atg ttg aac ttt gga tca gtc tct caa gga Ala Ser Ala Pro Ala Asn Met Leu Asn Phe Gly Ser Val Ser Gln Gly 320 325 330	1189
cca tcg agc gag tca tca gaa gag aat gag agc ggt tct cct gca atg Pro Ser Ser Glu Ser Ser Glu Glu Asn Glu Ser Gly Ser Pro Ala Met 335 340 345	1237
cac cgt gac aat aat aat ggg ata tat gga gct caa caa caa caa caa His Arg Asp Asn Asn Asn Gly Ile Tyr Gly Ala Gln Gln Gln Gln Gln 350 355 360 365	1285
caa caa cct ctt cat cct cat cag atg caa atg tac caa cat ctt tgg Gln Gln Pro Leu His Pro His Gln Met Gln Met Tyr Gln His Leu Trp 370 375 380	1333
tct aat cat ggt caa taa aatgaagcgg aaattaattt gtttccgttt Ser Asn His Gly Gln 385	1381
tggttacggt tatgggttga tttctt	1407
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<212> PRT	
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Pro Gly Met Leu Met Ser His Tyr Arg Asn Pro Asn Ala Ala Ala Ser 20 25 30	

mbi19 Sequence Listing.ST25

Pro Leu Met Val Pro Thr Ser Thr Ser Gln Pro Ile Gln His Pro Arg
 35 40 45

Leu Pro Phe Gly Asn Gln Gln Gln Ser Gln Thr Phe His Gln Gln Gln
 50 55 60

Gln Gln Gln Met Asp Gln Lys Thr Leu Glu Ser Leu Gly Phe Gly Asp
 65 70 75 80

Gly Ser Pro Ser Ser Gln Pro Met Arg Phe Gly Ile Asp Asp Gln Asn
 85 90 95

Gln Gln Leu Gln Val Lys Lys Lys Arg Gly Arg Pro Arg Lys Tyr Thr
 100 105 110

Pro Asp Gly Ser Ile Ala Leu Gly Leu Ala Pro Thr Ser Pro Leu Leu
 115 120 125

Ser Ala Ala Ser Asn Ser Tyr Gly Glu Gly Gly Val Gly Asp Ser Gly
 130 135 140

Gly Asn Gly Asn Ser Val Asp Pro Pro Val Lys Arg Asn Arg Gly Arg
 145 150 155 160

Pro Pro Gly Ser Ser Lys Lys Gln Leu Asp Ala Leu Gly Gly Thr Ser
 165 170 175

Gly Val Gly Phe Thr Pro His Val Ile Glu Val Asn Thr Gly Glu Asp
 180 185 190

Ile Ala Ser Lys Val Met Ala Phe Ser Asp Gln Gly Ser Arg Thr Ile
 195 200 205

Cys Ile Leu Ser Ala Ser Gly Ala Val Ser Arg Val Met Leu Arg Gln
 210 215 220

Ala Ser His Ser Ser Gly Ile Val Thr Tyr Glu Gly Arg Phe Glu Ile
 225 230 235 240

Ile Thr Leu Ser Gly Ser Val Leu Asn Tyr Glu Val Asn Gly Ser Thr
 245 250 255

Asn Arg Ser Gly Asn Leu Ser Val Ala Leu Ala Gly Pro Asp Gly Gly
 260 265 270

Ile Val Gly Gly Ser Val Val Gly Asn Leu Val Ala Ala Thr Gln Val
 275 280 285

Gln Val Ile Val Gly Ser Phe Val Ala Glu Ala Lys Lys Pro Lys Gln
 290 295 300

Ser Ser Val Asn Ile Ala Arg Gly Gln Asn Pro Glu Pro Ala Ser Ala
 305 310 315 320

Pro Ala Asn Met Leu Asn Phe Gly Ser Val Ser Gln Gly Pro Ser Ser
 325 330 335

mbil9 Sequence Listing.ST25

Glu Ser Ser Glu Glu Asn Glu Ser Gly Ser Pro Ala Met His Arg Asp
 340 345 350

Asn Asn Asn Gly Ile Tyr Gly Ala Gln Gln Gln Gln Gln Gln Pro
 355 360 365

Leu His Pro His Gln Met Gln Met Tyr Gln His Leu Trp Ser Asn His
 370 375 380

Gly Gln
 385

<210> 67
 <211> 1020
 <212> DNA
 <213> Arabidopsis thaliana

<220>
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 <222> (1)..(1020)
 <223> G2657

<400> 67
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 ttc cat gct aga gat ttc caa tta cat ctt caa caa caa caa caa cat 96
 Phe His Ala Arg Asp Phe Gln Leu His Leu Gln Gln Gln Gln Gln His
 20 25 30
 caa caa caa cat caa caa caa caa caa caa cag ttc ttt ctc cac cat 144
 Gln Gln Gln His Gln Gln Gln Gln Gln Gln Phe Phe Leu His His
 35 40 45
 cat cag caa cca caa aga aac ctt gat caa gat cac gag cag caa gga 192
 His Gln Gln Pro Gln Arg Asn Leu Asp Gln Asp His Glu Gln Gln Gly
 50 55 60
 ggg tca ata ttg aat aga tct atc aag atg gat cgc gaa gag aca agc 240
 Gly Ser Ile Leu Asn Arg Ser Ile Lys Met Asp Arg Glu Glu Thr Ser
 65 70 75 80
 gat aac atg gac aac atc gct aat acc aac agc ggt agc gaa ggt aaa 288
 Asp Asn Met Asp Asn Ile Ala Asn Thr Asn Ser Gly Ser Glu Gly Lys
 85 90 95
 gag atg agt tta cac gga gga gaa gga gga agc ggt ggt gga gga agt 336
 Glu Met Ser Leu His Gly Gly Glu Gly Gly Ser Gly Gly Gly Gly Ser
 100 105 110
 gga gaa cag atg aca aga agg cca aga gga aga cca gca gga tcc aag 384
 Gly Glu Gln Met Thr Arg Arg Pro Arg Gly Arg Pro Ala Gly Ser Lys
 115 120 125
 aac aaa cct aaa gct cca ata atc ata aca aga gac agc gca aac gcg 432
 Asn Lys Pro Lys Ala Pro Ile Ile Ile Thr Arg Asp Ser Ala Asn Ala
 130 135 140
 ctt cga act cac gtc atg gag ata gga gac gga tgt gac ata gtt gac 480
 Leu Arg Thr His Val Met Glu Ile Gly Asp Gly Cys Asp Ile Val Asp
 145 150 155 160
 tgt atg gct acg ttc gct aga cgc cgc caa aga ggc gtt tgc gtt atg 528
 Cys Met Ala Thr Phe Ala Arg Arg Arg Gln Arg Gly Val Cys Val Met
 165 170 175
 agc ggt aca gga agc gtt act aac gtc act ata cgt cag cct gga tcg 576
 Ser Gly Thr Gly Ser Val Thr Asn Val Thr Ile Arg Gln Pro Gly Ser

mbil9 Sequence Listing.ST25

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195	200	205	
ctt tgc gga tct ttc ttg cct ccg cct gcg ccg cct gca gcc acc gga	Leu Ser Gly Ser Phe Leu Pro Pro Pro Ala Pro Pro Ala Ala Thr Gly	672	
210	215	220	
cta agc gtt tac cta gcc gga gga caa ggg cag gtc gtt gga ggt agt	Leu Ser Val Tyr Leu Ala Gly Gly Gln Gly Gln Val Val Gly Gly Ser	720	
225	230	235	240
gtg gtg gga cct ttg ttg tgt tgc ggt cct gtg gtg gtt atg gcg gct	Val Val Gly Pro Leu Leu Cys Ser Gly Pro Val Val Val Met Ala Ala	768	
245	250	255	
tct ttt agc aat gcg gcg tac gaa agg ctg cct ttg gaa gaa gat gag	Ser Phe Ser Asn Ala Ala Tyr Glu Arg Leu Pro Leu Glu Glu Asp Glu	816	
260	265	270	
atg cag acg cca gtt caa gga ggc ggt gga gga gga gga ggt ggt ggt	Met Gln Thr Pro Val Gln Gly Gly Gly Gly Gly Gly Gly Gly Gly	864	
275	280	285	
gga atg gga tct ccc ccg atg atg gga cag caa caa gct atg gca gct	Gly Met Gly Ser Pro Pro Met Met Gly Gln Gln Gln Ala Met Ala Ala	912	
290	295	300	
atg gcg gcg gct caa gga cta cca ccg aat ctt ctt ggt tgc gtt cag	Met Ala Ala Ala Gln Gly Leu Pro Pro Asn Leu Leu Gly Ser Val Gln	960	
305	310	315	320
ttg cca ccg cca caa cag aat gat cag cag tat tgg tct acg ggt cgg	Leu Pro Pro Pro Gln Gln Asn Asp Gln Gln Tyr Trp Ser Thr Gly Arg	1008	
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cca ccg tat tga			1020
Pro Pro Tyr			

<210> 68
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 <212> PRT
 <213> Arabidopsis thaliana
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Phe His Ala Arg Asp Phe Gln Leu His Leu Gln Gln Gln Gln Gln His	20	25	30	
Gln Gln Gln His Gln Gln Gln Gln Gln Gln Phe Phe Leu His His	35	40	45	
His Gln Gln Pro Gln Arg Asn Leu Asp Gln Asp His Glu Gln Gln Gly	50	55	60	
Gly Ser Ile Leu Asn Arg Ser Ile Lys Met Asp Arg Glu Glu Thr Ser	65	70	75	80
Asp Asn Met Asp Asn Ile Ala Asn Thr Asn Ser Gly Ser Glu Gly Lys	85	90	95	
Glu Met Ser Leu His Gly Gly Glu Gly Gly Ser Gly Gly Gly Gly Ser				

mbi19 Sequence Listing.ST25

100

105

110

Gly Glu Gln Met Thr Arg Arg Pro Arg Gly Arg Pro Ala Gly Ser Lys
 115 120 125

Asn Lys Pro Lys Ala Pro Ile Ile Ile Thr Arg Asp Ser Ala Asn Ala
 130 135 140

Leu Arg Thr His Val Met Glu Ile Gly Asp Gly Cys Asp Ile Val Asp
 145 150 155 160

Cys Met Ala Thr Phe Ala Arg Arg Arg Gln Arg Gly Val Cys Val Met
 165 170 175

Ser Gly Thr Gly Ser Val Thr Asn Val Thr Ile Arg Gln Pro Gly Ser
 180 185 190

Pro Pro Gly Ser Val Val Ser Leu His Gly Arg Phe Glu Ile Leu Ser
 195 200 205

Leu Ser Gly Ser Phe Leu Pro Pro Pro Ala Pro Pro Ala Ala Thr Gly
 210 215 220

Leu Ser Val Tyr Leu Ala Gly Gly Gln Gly Gln Val Val Gly Gly Ser
 225 230 235 240

Val Val Gly Pro Leu Leu Cys Ser Gly Pro Val Val Val Met Ala Ala
 245 250 255

Ser Phe Ser Asn Ala Ala Tyr Glu Arg Leu Pro Leu Glu Glu Asp Glu
 260 265 270

Met Gln Thr Pro Val Gln Gly Gly Gly Gly Gly Gly Gly Gly Gly
 275 280 285

Gly Met Gly Ser Pro Pro Met Met Gly Gln Gln Gln Ala Met Ala Ala
 290 295 300

Met Ala Ala Ala Gln Gly Leu Pro Pro Asn Leu Leu Gly Ser Val Gln
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Pro Pro Tyr

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<400> 69

mbi19 Sequence Listing.ST25

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gatcttattc tccactgtat aaaatcagcg agattttaag ggattgtgaa ggtaccatct	180
taaacacaaa atg ggt act tct act aca gag agt gtg gtg gcg tgt gaa	229
Met Gly Thr Ser Thr Thr Glu Ser Val Val Ala Cys Glu	
1 5 10	
ttt tgc ggc gag aga acg gcg gtt ctg ttt tgt aga gcc gat acg gcg	277
Phe Cys Gly Glu Arg Thr Ala Val Leu Phe Cys Arg Ala Asp Thr Ala	
15 20 25	
aag ctt tgt ttg cct tgt gac cag cac gtg cac tcg gcg aac ctt ctc	325
Lys Leu Cys Leu Pro Cys Asp Gln His Val His Ser Ala Asn Leu Leu	
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tcg agg aag cat gtt cgt tct cag atc tgt gat aac tgt agc aaa gag	373
Ser Arg Lys His Val Arg Ser Gln Ile Cys Asp Asn Cys Ser Lys Glu	
50 55 60	
ccg gtg tcc gta cgt tgc ttc aca gat aat ctc gta ttg tgt cag gag	421
Pro Val Ser Val Arg Cys Phe Thr Asp Asn Leu Val Leu Cys Gln Glu	
65 70 75	
tgt gat tgg gat gtt cac gga agc tgt tcc tcc tcc gcg acg cat gaa	469
Cys Asp Trp Asp Val His Gly Ser Cys Ser Ser Ser Ala Thr His Glu	
80 85 90	
cgc tcc gcc gtg gaa ggg ttt tca ggt tgt cct tcg gtt ttg gag ctt	517
Arg Ser Ala Val Glu Gly Phe Ser Gly Cys Pro Ser Val Leu Glu Leu	
95 100 105	
gct gct gtg tgg gga atc gat tta aag ggt aag aag aaa gaa gat gac	565
Ala Ala Val Trp Gly Ile Asp Leu Lys Gly Lys Lys Lys Glu Asp Asp	
110 115 120 125	
gaa gac gaa ttg act aag aat ttt ggg atg ggg ttg gat tcg tgg ggt	613
Glu Asp Glu Leu Thr Lys Asn Phe Gly Met Gly Leu Asp Ser Trp Gly	
130 135 140	
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Ser Gly Ser Asn Ile Val Gln Glu Leu Ile Val Pro Tyr Asp Val Ser	
145 150 155	
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Cys Lys Lys Gln Ser Phe Ser Phe Gly Arg Ser Lys Gln Val Val Phe	
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Glu Gln Leu Glu Leu Leu Lys Arg Gly Phe Val Glu Gly Glu Gly Glu	
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Ile Met Val Pro Glu Gly Ile Asn Gly Gly Gly Ser Ile Ser Gln Pro	
190 195 200 205	
tct ccg acg acg tcg ttt act tct ttg ctt atg tct caa agt ctt tgt	853
Ser Pro Thr Thr Ser Phe Thr Ser Leu Leu Met Ser Gln Ser Leu Cys	
210 215 220	
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Gly Asn Gly Met Gln Trp Asn Ala Thr Asn His Ser Thr Gly Gln Asn	
225 230 235	
act cag ata tgg gat ttt aac ttg gga cag tcg agg aac cct gat gaa	949
Thr Gln Ile Trp Asp Phe Asn Leu Gly Gln Ser Arg Asn Pro Asp Glu	
240 245 250	
cct agt cca gtc gaa act aaa ggc tct act ttc aca ttc aac aac gtt	997
Pro Ser Pro Val Glu Thr Lys Gly Ser Thr Phe Thr Phe Asn Asn Val	
255 260 265	
act cat ctc aag aac gat acc cga acc acc aat atg aat gct ttc aaa	1045

mbi19 Sequence Listing.ST25

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          290          295          300

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Glu Thr Ser Lys Ser Asn Asn Ile Pro Ala Ala Ile His Ser His Lys
          305          310          315

agt tct aac gac tcc tgt ggc ttg cat tgc acg gaa cat att gct att   1189
Ser Ser Asn Asp Ser Cys Gly Leu His Cys Thr Glu His Ile Ala Ile
          320          325          330

act agt aat aga gcc aca aga ttg gtg gcg gta acg aat gct gat cta   1237
Thr Ser Asn Arg Ala Thr Arg Leu Val Ala Val Thr Asn Ala Asp Leu
          335          340          345

gag cag atg gca cag aac aga gat aat gct atg cag cgg tac aag gaa   1285
Glu Gln Met Ala Gln Asn Arg Asp Asn Ala Met Gln Arg Tyr Lys Glu
          350          355          360          365

aag aag aaa acg cgg aga tat gat aag acc ata aga tat gaa acg agg   1333
Lys Lys Lys Thr Arg Arg Tyr Asp Lys Thr Ile Arg Tyr Glu Thr Arg
          370          375          380

aag gcg aga gcc gag acc agg ttg cgt gtt aag ggc aga ttt gtg aaa   1381
Lys Ala Arg Ala Glu Thr Arg Leu Arg Val Lys Gly Arg Phe Val Lys
          385          390          395

gct aca gat cct tag atgtctctcc acgttaggtt ttacatttga gacctaagt   1436
Ala Thr Asp Pro
          400

taggaacttt ttttgttttt tctactttca actaccttgt aaatgtaaat gatcgatctt   1496

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<210> 70
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<400> 70

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Glu Arg Thr Ala Val Leu Phe Cys Arg Ala Asp Thr Ala Lys Leu Cys
          20          25          30

Leu Pro Cys Asp Gln His Val His Ser Ala Asn Leu Leu Ser Arg Lys
          35          40          45

His Val Arg Ser Gln Ile Cys Asp Asn Cys Ser Lys Glu Pro Val Ser
          50          55          60

Val Arg Cys Phe Thr Asp Asn Leu Val Leu Cys Gln Glu Cys Asp Trp
65          70          75          80

Asp Val His Gly Ser Cys Ser Ser Ser Ala Thr His Glu Arg Ser Ala
          85          90          95

Val Glu Gly Phe Ser Gly Cys Pro Ser Val Leu Glu Leu Ala Ala Val
          100          105          110

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mbi19 Sequence Listing.ST25

Trp Gly Ile Asp Leu Lys Gly Lys Lys Lys Glu Asp Asp Glu Asp Glu
 115 120 125
 Leu Thr Lys Asn Phe Gly Met Gly Leu Asp Ser Trp Gly Ser Gly Ser
 130 135 140
 Asn Ile Val Gln Glu Leu Ile Val Pro Tyr Asp Val Ser Cys Lys Lys
 145 150 155 160
 Gln Ser Phe Ser Phe Gly Arg Ser Lys Gln Val Val Phe Glu Gln Leu
 165 170 175
 Glu Leu Leu Lys Arg Gly Phe Val Glu Gly Glu Gly Glu Ile Met Val
 180 185 190
 Pro Glu Gly Ile Asn Gly Gly Gly Ser Ile Ser Gln Pro Ser Pro Thr
 195 200 205
 Thr Ser Phe Thr Ser Leu Leu Met Ser Gln Ser Leu Cys Gly Asn Gly
 210 215 220
 Met Gln Trp Asn Ala Thr Asn His Ser Thr Gly Gln Asn Thr Gln Ile
 225 230 235 240
 Trp Asp Phe Asn Leu Gly Gln Ser Arg Asn Pro Asp Glu Pro Ser Pro
 245 250 255
 Val Glu Thr Lys Gly Ser Thr Phe Thr Phe Asn Asn Val Thr His Leu
 260 265 270
 Lys Asn Asp Thr Arg Thr Thr Asn Met Asn Ala Phe Lys Glu Ser Tyr
 275 280 285
 Gln Glu Asp Ser Val His Ser Thr Ser Thr Lys Gly Gln Glu Thr Ser
 290 295 300
 Lys Ser Asn Asn Ile Pro Ala Ala Ile His Ser His Lys Ser Ser Asn
 305 310 315 320
 Asp Ser Cys Gly Leu His Cys Thr Glu His Ile Ala Ile Thr Ser Asn
 325 330 335
 Arg Ala Thr Arg Leu Val Ala Val Thr Asn Ala Asp Leu Glu Gln Met
 340 345 350
 Ala Gln Asn Arg Asp Asn Ala Met Gln Arg Tyr Lys Glu Lys Lys Lys
 355 360 365
 Thr Arg Arg Tyr Asp Lys Thr Ile Arg Tyr Glu Thr Arg Lys Ala Arg
 370 375 380
 Ala Glu Thr Arg Leu Arg Val Lys Gly Arg Phe Val Lys Ala Thr Asp
 385 390 395 400

Pro

INTERNATIONAL SEARCH REPORT

International application No.

PCT/US00/31414

A. CLASSIFICATION OF SUBJECT MATTER

IPC(7) : A01H 1/00, 5/00; C12N 5/04, 15/00, 15/82; C12P 21/02
US CL : 435/69.1, 320.1, 410, 419, 468; 800/278, 284, 287, 290

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)
U.S. : 435/69.1, 320.1, 410, 419, 468; 800/278, 284, 287, 290

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)
East, USPAT; STN, Agricola, Biosis, CaPlus, Embase; Sequence Search of SEQ ID NOs. 1 & 2

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	AOYAMA, T. et al. Ectopic expression of the Arabidopsis transcriptional activator Athb-1 alters leaf cell fate in tobacco, The Plant Cell, November 1995, Vol. 7, pages 1773-1785, entire document.	1-10, 13, 25
---		-----
Y		11, 12, 26
X	BELLIS, L.D. et al. Distinct cis-acting sequences are required for the germination and sugar responses of the cucumber isocitrate lyase gene. Gene 1997, Vol. 197, pages 375-378, entire document.	1-10, 13, 25
X	KIM, S. et al. Sugar response element enhances wound response of potato proteinase inhibitor II promoter in transgenic tobacco. Plant Mol. Biol. 1991, Vol. 17, pages 973-983, entire document.	1-10, 13, 25
Y	Database Genbank on NCBI, US National Library of Medicine, (Bethesda, MD, USA), No. U78721, LIN, X. et al. 'Sequence and analysis of chromosome 2 of the plant Arabidopsis thaliana' 5 April 2000, especially bases 14,116-14,895.	1-10, 13, 25

☐

Further documents are listed in the continuation of Box C.

☐

See patent family annex.

* Special categories of cited documents:		"T"	later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
"A"	document defining the general state of the art which is not considered to be of particular relevance	"X"	document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
"E"	earlier application or patent published on or after the international filing date	"Y"	document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art
"L"	document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)	"&"	document member of the same patent family
"O"	document referring to an oral disclosure, use, exhibition or other means		
"P"	document published prior to the international filing date but later than the priority date claimed		

Date of the actual completion of the international search

23 February 2001 (23.02.2001)

Date of mailing of the international search report

04 APR 2001

Name and mailing address of the ISA/US

Commissioner of Patents and Trademarks
Box PCT
Washington, D.C. 20231

Facsimile No. (703)305-3230

Authorized officer

David Kruse

Telephone No. 703-308-1600

TERRY J. DEY
PARALEGAL SPECIALIST
TECHNOLOGY CENTER 1600

INTERNATIONAL SEARCH REPORT

International application No.

PCT/US00/31414

Box I Observations where certain claims were found unsearchable (Continuation of Item 1 of first sheet)

This international report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☐ Claim Nos.:
because they relate to subject matter not required to be searched by this Authority, namely:
2. ☐ Claim Nos.:
because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out, specifically:
3. ☒ Claim Nos.: 14 & 24
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box II Observations where unity of invention is lacking (Continuation of Item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:
Please See Continuation Sheet

1. ☐ As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims.
2. ☐ As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3. ☐ As only some of the required additional search fees were timely paid by the applicant, this international search report covers only those claims for which fees were paid, specifically claims Nos.:
4. ☒ No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.: 1-13, 25 and 26; SEQ ID NOs 1 & 2

Remark on Protest

☐
☐

The additional search fees were accompanied by the applicant's protest.

No protest accompanied the payment of additional search fees.

BOX II. OBSERVATIONS WHERE UNITY OF INVENTION IS LACKING This application contains the following inventions or groups of inventions which are not so linked as to form a single general inventive concept under PCT Rule 13.1. In order for all inventions to be examined, the appropriate additional examination fees must be paid.

Groups I-XXXV, claim(s) 1-13 and 25-26, drawn to a transgenic plant having modified seed characteristics, polynucleotides and vectors for producing said transgenic plant and a method of making said transgenic plant. Applicant must elect one pair of sequences (one nucleic acid and the corresponding amino acid translation) to be examined, *i.e.* SEQ ID NO: 1 and 2 in Group I, SEQ ID NO: 3 and 4 in Group II, SEQ ID NO: 5 and 6 in Group III, etc.

Group XXXVI, claim(s) 15-17, drawn to a method of identifying a factor that is modulated.

Group XXXVII, claims(s) 18, drawn to a method of identifying a molecule that modulates activity or expression of a polynucleotide or polypeptide.

Group XXXVIII, claims(s) 19 and 20, drawn to an integrated computer system.

Group XXXIV, claim(s) 21-23, drawn to a method for identifying a polynucleotide sequence comprising selecting a nucleic acid sequence from a database that meets a selected sequence criteria.

The inventions listed as Groups I-XXXIV do not relate to a single general inventive concept under PCT Rule 13.1 because, under PCT Rule 13.2, they lack the same or corresponding special technical features for the following reasons:

The inventions listed as Groups I-XXXIX do not relate to a single general inventive concept under PCT Rule 13.1 because, under PCT Rule 13.2, they lack the same or corresponding special technical features for the following reasons: Groups I-XXXV are drawn to a transgenic plant and a method of producing said plant with a nucleic acid sequence. The methods of Groups I-XXXV differ from each other in that they are directed to a plant transformation method and transgenic plant with a structurally and functionally distinct nucleic acid sequence which encodes a structurally and functionally distinct amino acid sequence. In addition, Groups XXXVI, XXXVII and XXXIX are different methods from any of Groups I-XXXV in that they have different method steps and different end products, and Group XXXVIII requires a computer system. Thus, there is no single special technical feature, which links the inventions of Groups I-XXXIX under PCT Rule 13.2.